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(21) International Application Number: PCT/US92/01300 (22) International Filing Date: 20 February 1992 (20.02.92) (30) Priority data: 657,236 22 February 1991 (22.02.91) US (60) Parent Application or Grant (63) Related by Continuation US 657,236 (CIP) Filed on 22 February 1991 (22.02.91) (71) Applicant (for all designated States except US): AMERICAN CYANAMID COMPANY [US/US]; 1937 West Main Street, P.O. Box 60, Stamford, CT 06904 (US).		(72) Inventors; and (75) Inventors/Applicants (for US only) : Terman, Bruce, Israel [US/US]; 17 Prospect Street, Monroe, NY 10950 (US). CARRION, Miguel, Eduardo [EC/US]; 26 Summit Avenue, Spring Valley, NY 10977 (US). (74) Agent: GORDON, Alan, M.; American Cyanamid Company, 1937 West Main Street, P.O. Box 60, Stamford, CT 06904 (US). (81) Designated States: AT (European patent), BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, KR, LU (European patent), MC (European patent), NL (European patent), SE (European patent), US. Published <i>With international search report.</i>
(54) Title: IDENTIFICATION OF A NOVEL HUMAN RECEPTOR TYROSINE KINASE GENE (57) Abstract A DNA sequence encoding a novel human growth factor receptor referred to as a type III receptor tyrosine kinase is described. The amino acid sequence of the receptor is also described. The receptor has a sequence which is similar to that of the kinase domains of known type III receptor tyrosine kinases, but which is unique in its kinase insert domain sequence. The receptor binds specifically to the vascular endothelial cell growth factor.		

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IDENTIFICATION OF A NOVEL HUMAN
RECEPTOR TYROSINE KINASE GENE

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FIELD OF THE INVENTION

This invention relates to the DNA sequence encoding a novel human growth factor receptor which is a type III receptor tyrosine kinase. The receptor is referred to as Kinase insert Domain containing Receptor (KDR) and binds specifically to the growth factor vascular endothelial cell growth factor (VEGF). This invention also relates to the amino acid sequence of the receptor.

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BACKGROUND OF THE INVENTION

Growth factors are small molecules which regulate normal cell growth and development through interaction with cell surface receptors. The receptors for a number of growth factors are referred to as tyrosine kinases; that is, binding of growth factor to the receptor stimulates an increased phosphorylation of tyrosine amino acids within the receptor; this in turn leads to cellular activation (Bibliography 1).

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There is increasing evidence that genetic alterations affecting the expression of receptor tyrosine kinases (RTK) can contribute to the altered cell growth associated with cancer. This conclusion is

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supported by the frequent identification of RTK as products of the oncogenes for many of the acutely transforming retroviruses (e.g., 2,3,4) and the overexpression of RTK in certain cancers (5). The identification of a novel RTK may lead to a better understanding of cell growth under both normal and transforming circumstances.

The amino acid sequence in the catalytic domain of all tyrosine kinases has been conserved (6). Detailed analysis of the amino acid sequences within the catalytic and noncatalytic domains of RTK indicates the existence of distinct structural subtypes. One group of RTK (designated type III) includes the ckit proto-oncogene and the receptors for platelet derived growth factor (PDGF) and colony stimulating factor-1 (CSF-1).

The most unusual feature of this subtype is that its catalytic (kinase) domain is interrupted by a long insertion sequence of 12-102 amino acids (the kinase insert domain). The two peptides constituting the kinase domain are conserved between the receptors, while the sequence of the kinase insert domain is unique for each receptor.

Several approaches have been tried in order to identify novel RTK, including low-stringency screening of cDNA libraries with previously characterized DNA probes (7). More recently, a technique has been developed that is capable of greatly facilitating the identification of novel genes for which some sequence data are known. The polymerase chain reaction (PCR) has been used to identify novel members of several gene families including those of guanine nucleotide regulatory proteins (8) and protein phosphatases (9). PCR has been used to identify novel tyrosine kinase genes (10), though the primers used in

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that study were designed from DNA segments contained in all tyrosine kinases, rather than being specifically directed against RTK. It is a continuing goal to identify receptors for growth factors.

5 The elucidation of the growth factors, as well as their receptors, involved in regulating endothelial cell function is critical for the understanding of how new blood vessels are formed (angiogenesis). Angiogenesis plays a significant role
10 in both normal and pathological events such as embryogenesis, progression of ocular diseases, and wound healing (11). In particular, angiogenesis is an important process for the growth of tumors (11). Angiogenesis is a complex process involving endothelial
15 cell proliferation, migration, and tissue infiltration. These events are stimulated by growth factors which either (i) act directly on endothelial cells (12,13), or (ii) act indirectly by inducing host cells to release specific endothelial cell growth factors (11).
20 One member of the first group is vascular endothelial cell growth factor (VEGF), also known as vascular permeability factor (14-16). Besides its angiogenic activity, VEGF displays the physiological function of increasing the permeability of capillary vessels to
25 different macromolecules (14).

SUMMARY OF THE INVENTION

30 The present invention relates to novel DNA segments which together comprise a gene which encodes type III RTK. The type III RTK encoded by the gene is designated the KDR protein (which stands for Kinase insert Domain containing Receptor). The KDR protein
35 binds specifically to the growth factor VEGF (vascular endothelial cell growth factor).

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The DNA segments are identified and isolated through the use of PCR technology. The overall strategy is summarized as follows:

5 PCR is used to amplify the DNA segments corresponding to the kinase insert domains of type III receptor tyrosine kinase genes in an endothelial cell library designated HL10246 (Clontech Laboratories, Inc., Palo Alto, CA). Degenerate oligonucleotide primers are designed which are complementary to
10 conserved tyrosine kinase domains flanking the kinase insert domains of known type III receptor tyrosine kinases. These primers are used in the PCR procedure. DNA probes, designed from the DNA sequence of the PCR product, are then used to identify cDNA clones of the
15 receptor gene from the original cDNA library.

In particular, the present invention relates to specific oligonucleotides which, when used as primers for PCR, allow for the amplification of DNA segments corresponding to the kinase insert domains of
20 type III RTK genes.

In a principal embodiment, the present invention is directed to three overlapping DNA segments (designated BTIII1081.8, BTIII129.5 and BTIV169) which
25 comprise the entire coding region of this novel gene, namely, 4,068 nucleotides extending to the 3' end.

These DNA segments are isolated from a human endothelial cell cDNA library and together comprise the gene coding for a novel type III receptor tyrosine kinase. The human gene containing these DNA segments
30 is referred to hereinafter as KDR (which stands for Kinase insert Domain containing Receptor) or, alternatively, as kdp (which stands for Kinase insert Domain containing Protein). The use of the term KDR is intended to include any DNA segments which form the
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human gene which encodes the novel type III RTK of this application.

The DNA segments embodied in this invention are isolated from human sources. The present invention comprises DNA segments, and methods for using these DNA segments, which allow for the identification of a closely related gene in mouse DNA. The methods developed in this invention can be readily used by those skilled in the art for the identification and isolation of closely-related homologues in other species. Therefore, the present invention also embodies all DNA segments from species other than human which encode proteins having substantially the same amino acid sequence as that encoded by the kdp gene.

The present invention further relates to methods developed for the detection of mRNA's produced as a result of transcription of the sense strands of the DNA segments of this invention. Messenger RNA prepared from bovine endothelial cells are used in developing these methods. The ability to detect mRNA for a novel RTK may ultimately have medical benefit, especially in light of recent observations that the mRNA for certain RTKs are overexpressed in some cancers (5).

The methods developed in the present invention for detecting mRNA expressed by the kdp gene can be readily used by those of ordinary skill in the art for the detection of mRNA species related to the kdp gene in any cell type and from any species. For this reason, the present invention embodies all mRNA segments which are the result of transcription of the kdp gene.

The present invention relates to methods for expression of the receptor protein, for example, in CMT-3 cells of monkey kidney origin. The receptor

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protein, portions thereof, and mutated forms of the receptor protein may be expressed in many other cells by those skilled in the art using methods similar to those described in this application. For this reason, the present invention embodies all proteins encoded by the human KDR gene and proteins encoded by related genes found in other species.

The present invention further relates to methods for studying the interaction of VEGF to the expressed KDR protein. Recent work in the literature (17) indicates that VEGF is one member of a family of related proteins, and the interaction of growth factors similar to VEGF with the KDR protein can readily be studied by those skilled in the art using methods similar to those described in this application. These methods can readily be modified to study the interaction of candidate pharmaceuticals with the KDR protein towards the goal of developing an antagonist or agonist of VEGF action. For this reason, the present invention embodies methods for studying the interaction of VEGF and VEGF-related growth factors with the KDR protein.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts a schematic representation of three receptor tyrosine kinase subclasses (6). KI is kinase insert domain; PTK is kinase domain; cys is cysteine rich region.

Figure 2 depicts the two sets of primers used for PCR (SEQ ID No: 1 and 2). The nucleotide sequences in appropriate regions of the four known type III receptor tyrosine kinase cDNAs are aligned and degenerate oligonucleotide primers are designed based upon the consensus sequences.

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Figure 3 depicts the amplification of the kinase insert domains using PCR. DNA segments encoding the kinase insert domains of type III receptor tyrosine kinases are amplified by PCR. A sample (5 μ l) is run on a 1.0% agarose gel which is stained with ethidium bromide. DNA size standards (123 bp ladder; Bethesda Research Laboratories, Bethesda, MD) are run as well.

Figure 4 depicts the DNA sequence of the two PCR products (Panel A: 363 bp segment derived from the 420 bp product (SEQ ID NO: 3); Panel B: 251 bp product (SEQ ID NO: 4)). The two products are purified by agarose gel electrophoresis, digested with Sall and EcoRI, and cloned into the plasmid vector pBlueScribe(+) (Stratagene; San Diego, CA). The 420 bp PCR product is digested to 363 bp during this procedure. The DNA sequences for the primers used in the amplification are underlined.

Figure 5A depicts a computer assisted comparison of the DNA sequence for the 363 bp DNA segment derived from the 420 bp PCR product with the sequence of a DNA segment of the PDGF receptor (SEQ ID NO: 5) (18). A region of strong homology between the 363 bp segment derived from the 420 bp PCR product and the PDGF receptor is contained in a box. Figure 5B depicts a computer assisted comparison of the DNA sequence for the 251 bp PCR product with the sequence of a DNA segment of the FGF receptor (SEQ ID NO: 6) (7).

Figure 6 depicts the strategy used for sequencing the insert portions of clones BTIII081.8 and BTIII129.5 and BTIV169. The sequencing reaction uses either synthetic oligonucleotides (represented by boxes at the start of an arrow), or the M13 universal primer (no box) to initiate the reaction. In some cases, portions of these DNA segments are isolated using the

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restriction enzymes indicated in the figure, and subcloned back into the plasmid vector pUC118, so that the M13 universal primer can be used. The position of the stop codon in BTIII129.5 is indicated. The coding portions of these DNA segments are shown at the bottom of the figure. The relative positions of the 1) membrane spanning portion, 2) kinase domains, and 3) kinase insert domain are indicated. The position of these structural features within the KDR derived DNA segments is compared in relation to their position in the PDGF-receptor ("PDGF-R").

Figure 7 depicts the DNA and predicted amino acid sequence of KDR, plus the stop codon (nucleotides 1-4071 of SEQ ID NO. 7). The sequence of the DNA segment amplified by PCR is underlined (nucleotides 2749-3105 of SEQ ID NO. 7). Cysteine residues in the putative extracellular domain are circled. Potential N-linked glycosylation sites are indicated by an asterisk. The putative membrane spanning region is enclosed in a box (nucleotides 2293-2367 of SEQ ID NO. 7).

Figure 8 depicts a hydropathy plot of the predicted amino acid sequence for the KDR protein.

Figure 9 depicts a comparison of the predicted amino acid sequence in the putative intracellular portion of the KDR protein to the ckit proto-oncogene (SEQ ID No: 8) (3), the CSF-1 receptor (SEQ ID NO: 9) (4), and the PDGF receptor (SEQ ID NO: 10) (18). Exact matches are indicated by an asterisk. Gaps are introduced to achieve maximum alignment. The putative ATP recognition site is indicated by three asterisks.

Figure 10 depicts the identification of kdp receptor mRNA by Northern blot analysis. Five micrograms of bovine aortic endothelial cell polyA+ RNA

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are used. A nick-translated [32 P] CTP-labelled EcoRI/BamHI DNA segment (nucleotides 1510-2417 of SEQ ID NO. 7) is used as a probe. Autoradiography is for 36 hours.

5 Figure 11 depicts the kdp gene in human and mouse DNA by Southern blot analysis. A nick translated [32 P]CTP-labelled EcoRI/BamHI DNA segment (nucleotides 1510-2417 of SEQ ID NO. 7) is used as the probe. The probe is hybridized to Southern blots containing EcoRI digested DNA from human (lane 1), mouse (lane 2), and human-mouse hybrid cells (19) (lanes 3 and 4). The DNA used in lane 3 lacks the kdp locus, while DNA used in lane 4 contains the kdp locus.

10 Figure 12 depicts a Western blot analysis of CMT-3 cells which express the KDR protein. Cells are transfected with either the pCDNA1tkpASP vector alone (lane 1) or with that vector modified to contain the KDR gene (lane 2). 2×10^5 cells and 1 microgram of DNA are used for each transfection. Forty-eight hours later, Western blot analysis is performed on the samples using the anti-KDR.PS23 polyclonal antibody at a dilution of 1:1000. Detection of reacting proteins is performed using an ECL system (Amersham, Chicago, IL).

15 Figure 13 depicts the results of [125 I] VEGF binding to CMT-3 cells which express the KDR protein. Cells are transfected with either the vector alone (bars 1 and 2) or with the vector containing the KDR gene (bars 3 and 4). Forty-eight hours later, the samples are washed with phosphate buffered saline (PBS), and incubated with serum-free media containing 50 pM [125 I] VEGF (specific activity equal to 4,000 cpm per fmol), for 90 minutes. Nonradioactive VEGF, 5 nM, is added to some samples (bars 2 and 4) to define

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specific binding sites. The samples are washed with ice cold PBS, and the cells are transferred to gamma-counting tubes using 0.1% lubrol.

Figure 14 depicts the results of affinity cross-linking of [¹²⁵I] VEGF to CMT-3 cells which express the KDR protein. CMT-3 cells are transfected with either the vector alone (lane 1) or with the vector containing the KDR gene (lane 2). Forty-eight hours later, the cells are washed in PBS, and serum free media containing 200 pM [¹²⁵I] VEGF is added. After 90 minutes at room temperature, an affinity cross-linker disuccinimidyl suberate, 0.5 mM, is added for 15 minutes. The samples are then prepared for SDS-PAGE autoradiography.

DETAILED DESCRIPTION OF THE INVENTION

The strategy used to discover the DNA segments for the novel type III RTK gene begins with the design of two degenerate oligonucleotide primers based upon their homology to specific regions of the kinase domains of known RTK genes (Fig. 2) (3,4,7,18). In one embodiment, the polymerase chain reaction is then used to amplify DNA segments from a human endothelial cell cDNA library (designated HL 10246). The cDNA products from this step are each cloned into a plasmid vector designated pBlueScribe+ (Stratagene, San Diego, CA) and sequenced. Oligonucleotide probes are designed from potentially interesting sequences in order to screen the cDNA library for more full length clones of the novel cDNA.

The strategy just described provides several novel elements: 1) the DNA sequences of the oligonucleotide primers used during PCR; 2) the DNA sequence of the products generated by the polymerase chain

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reaction; and 3) the DNA sequence of the final cloned DNA segments. Each of these elements of the invention described in this application will now be discussed in detail.

5 Figure 2 shows the rationale for choosing the oligonucleotide primers used in the PCR. The primers are designed to allow for the PCR amplification of the kinase insert domain of type III RTK genes. In order to design the primers, the DNA sequences of known type III
10 RTK genes are aligned in specific regions of their catalytic domains, and a consensus sequence is chosen. The regions of the catalytic domains chosen in designing the primers flank the kinase insert domains of the receptor genes.

15 Primer 1 (SEQ ID No: 1) is designed from a region of the kinase domain 5' to the kinase insert domain, and consists of a mixture of four different 21mers. Primer 2 (SEQ ID NO: 2) is designed from a region of the kinase domain 3' to the kinase insert
20 domain, and consists of a mixture of sixteen different 29mers with one inosine, indicated in SEQ ID NO: 2 by "N".

25 SalI and EcoRI restriction sites are included at the 5' end of primers 1 and 2, respectively, to facilitate the subcloning of the amplified PCR products into plasmid vectors. Those skilled in the art may use other restriction sites; other minor modifications in the protocol above permits the design of primers without the inclusion of restriction sites.

30 The selection of these specific primers constitutes a novel approach towards identifying novel type III RTK genes. It had previously been shown (10) that primers designed from DNA sequences common to all tyrosine kinases allows for the identification of novel
35 proteins. The present invention is the first to

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contemplate the use of PCR to specifically target type III RTK.

The protocol used for PCR is as follows:
Human endothelial cell cDNA (designated HL10246) is
denatured by boiling and submitted to 30 cycles of PCR
using 1 nmol of both primers in a final volume of 100
 μ l. The timing is 1.5 minutes at 92°C, 2 minutes at
50°C, and 2 minutes at 74°C. DNA from 5 μ l of sample is
separated on a 1% agarose gel and stained with ethidium
bromide.

Figure 3 shows the results of the PCR
amplification. Two DNA products, with sizes 251 bp
(SEQ ID NO: 4) and 420 bp, are visible when a sample of
the reaction is electrophoresed on a 1.0% agarose gel
and stained with ethidium bromide. The sizes of the
two products are within the range expected for type III
RTK genes (products derived from the FGF and PDGF
receptor genes, which have the smallest and largest
known kinase insert domains, would be 230 and 510 bp,
respectively (20, 21).

The DNA from four contiguous lanes with
sizes ranging from 200 to 600 bp is electrophoresed
onto DEAE filter paper, eluted from the paper with
salt, and ethanol precipitated. The samples are
incubated with 5 units of EcoRI and SalI. The
restriction enzymes digest the 420 bp DNA segment to a
363 bp DNA segment (SEQ ID NO: 3), due to the presence
of an EcoRI site within the 420 bp DNA segment
(nucleotide 2749, SEQ ID NO. 7). The restriction
enzyme digested PCR products are then subcloned into
the plasmid vector pBlueScribe(+). The recombinant
clones are analyzed by sequencing using the
dideoxy-method (22) using a United States Biochemical
(Cleveland, Ohio) Sequenase Version 2.0 sequencing kit.
Figure 4 shows the DNA sequences for the 251 bp PCR

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product and the 363 bp DNA segment derived from the 420 bp PCR product.

Computer assisted comparison of the DNA sequence for the 363 bp segment of the 420 bp PCR product to databases of known DNA sequences reveals that the sequence is novel, because it shares strong sequence identity with the flanking catalytic domain of known type III RTK genes, but not their kinase insert domains. Figure 5A compares the DNA sequence for the 363 DNA segment with that for the PDGF receptor gene (SEQ ID No: 5). Similar results are obtained using other type III RTK genes.

DNA sequencing of the 251 bp PCR product reveals a novel sequence containing both primers used for the amplification, but the sequence shows little homology to known tyrosine kinases. This is depicted in Figure 5B, which compares the DNA sequence for the 251 bp DNA segment with that for the FGF receptor (SEQ ID NO: 6). For this reason, further analysis of Product 1 is not pursued.

The protocols used during the PCR do not allow for amplification of the kinase insert domains of known receptor tyrosine kinases in the endothelial cell library used because of the low copy number of the message present in the library. There have been many studies on the effect of FGF on endothelial cell function (23,24) although there is evidence that the expression of the FGF receptor is developmentally regulated (7) and it is likely that the library used contains little or no cDNA for the FGF receptor.

An oligonucleotide probe, designed from the DNA sequence of the 363 bp segment, is synthesized (using an ABI 380 DNA Synthesizer) in order to screen the human endothelial cell cDNA library (HL10246) for the isolation of more full length clones containing the

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363 bp DNA segment. The probe sequence is chosen from the region of the 363 bp DNA segment which shares little sequence homology with known RTK.

The screening of the endothelial cell cDNA library is conducted as follows: Lambda gt11 phage, 10⁶, are adsorbed to *E. coli* LE392 for 15 minutes at 37°C prior to plating onto agar plates at a density of 5 x 10⁵ phage per plate. After allowing the phage plaques to develop at 37°C, plaque lifts are made using nitrocellulose filters, denatured in 0.4 N NaCl for 1 minute, and neutralized in 0.5 M Tris.HCl, pH 7.3, plus 1.5 M NaCl. The filters are washed with 2 x standard saline citrate (SSC) and then baked for 1.5 hour in a vacuum oven at 80°C. The filters are probed with an [³²P] ATP end labeled synthetic oligonucleotide, 5' - TTTCCCTTGACGGAATCGTGCCCTTTGGT-3', which is the reverse complement of a DNA sequence contained in the PCR amplified product (Fig. 3). Hybridization is performed at 50°C in 5 x SSPE (167 mM NaCl, 10 mM sodium phosphate, pH 7.4, 1 mM EDTA), 2.5 x Denhardts, 0.5% sodium dodecyl sulfate (SDS), 100 µg/ml salmon sperm DNA. The filters are washed twice, 20 minutes per wash, with 2 x SSC plus 0.1% SDS at room temperature, followed by washing twice at 50°C with 0.1 X SSC plus 0.1% SDS; 20 minutes per wash. Positive clones are identified, picked and plaque purified.

Forty-five positive clones are obtained. Three of these positive clones are plaque purified and their phage DNA isolated. Digestion of the DNA with *Eco*RI and electrophoresis in agarose indicates that one clone, designated BTIII081.8, contains the largest insert, and subsequent analysis indicates that the DNA insert of this clone overlaps that of the inserts contained in other two purified clones (designated BTIII079.11 and BTIII079.47A).

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Digestion of the purified phage DNA of the clone designated BTIII081.8 with EcoRI results in DNA segments of 250 bp, 600 bp, and 1000 bp. Each of these three products is subcloned into the plasmid vector pUC118 and sequenced (Figure 6 shows the strategy used for sequencing). The orientation of the three fragments is determined by subcloning from the insert a BglII/BglII fragment into pUC118 and sequencing across the EcoRI junctions using a synthetic oligonucleotide to prime the sequencing reaction.

A restriction map is determined for each fragment (Figure 6). Various restriction site pieces are removed from the plasmids and recloned into pUC118 so that sequencing the resulting plasmids with the universal primer allows for sequencing most of the entire original fragments in both directions. Three oligonucleotide primers are required to sequence the entire cDNA in both directions. For the purposes of this application, this insert contains nucleotides numbered 1510-3406 (SEQ ID NO. 7).

A [³²P]CTP-labelled, nick-translated EcoRI-BamHI DNA segment derived from clone BTIII081.8 (nucleotides 1510-2417 of SEQ ID NO. 7) is used as a probe to rescreen the original endothelial cell cDNA library for more 5' full length DNA segments of the gene from which the insert portion of BTIII081.8 is derived. The protocols used to isolate the overlapping clones are identical to that used to isolate BTIII081.8.

A synthetic oligonucleotide probe is designed with 29 nucleotides corresponding to part of the DNA sequence of the insert portion of the clone BTIII081.8 (nucleotides 3297-3325 of SEQ ID NO. 7) in order to rescreen the original endothelial cell cDNA library for more full 3' length DNA segments of the gene from which

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the insert portion of BTIII081.8 is derived. The protocols used to isolate the overlapping clones are identical to that used to isolate BTIII081.8. Several positive clones for each of the 5' and 3' ends are identified and plaque purified.

One of the clones is designated BTIII200.2. The DNA from BTIII200.2 contains a 3.4 kb insert as determined by EcoRI digestion of the isolated phage DNA. EcoRI digestion of BTIII200.2 results in three DNA fragments. One of these fragments (2.5 kb) is cloned into pUC119 and is designated BTIV006. The clone BTIV006 contains nucleotides numbered 7-2482. As described below, BTIV006 plus nucleotides 1-6 is designated BTIV169. DNA sequencing of the 2.5 kb DNA insert (BTIV169) indicates that it overlaps over one thousand nucleotides of the DNA sequence of the insert portion of the clone BTIII081.8 (Figure 6) at the 5' end.

A second clone isolated from the cDNA library is designated BTIII129.5. The DNA from BTIII129.5 contains a 2.2 kb insert as determined by EcoRI digestion of the isolated phage DNA. DNA sequencing of the 2.2 kb DNA insert indicates that it overlaps over five hundred nucleotides of the DNA sequence of the insert portion of the clone BTIII081.8 (Figure 6). The clone BTIII129.5 contains nucleotides numbered 2848-4236 (SEQ ID NO. 7). The DNA sequence for BTIII129.5 contains the stop codon TAA, defining the position of the 3' end of an open reading frame for the novel gene. Except for the first six nucleotides of the gene which are discussed below, these three clones define a gene encoding a growth factor receptor. These three clones define a 4,062 nucleotide sequence of the open reading frame of the gene extending to the 3' end, followed by a 168 nucleotide non-coding region (SEQ ID

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NO. 7). A sample of a lambda gt11 phage harboring the clone BTIII081.8 has been deposited with the American Type Culture Collection, 12301 Parklawn Drive, Rockville, Maryland 20852, U.S.A., and has been assigned ATCC accession number 40,931. A sample of a lambda gt11 phage harboring the clone BTIII129.5 has been deposited with the American Type Culture Collection and has been assigned ATCC accession number 40,975. For reasons discussed below, a sample of the clone BTIV006 was not deposited.

The aforementioned DNA segments (BTIII081.8, BTIII129.5, and BTIII200.2 (or BTIV006) encode 4062 nucleotides of the coding portion of a novel gene. The cDNA clones are incomplete in that a transcription initiation coding for methionine is missing. After the isolation of these clones, Matthews et al. (25) reported the cloning of a gene homologue of KDR in mouse, which was referred to as Flk-1. Analysis of the nucleic acid and amino acid sequence of Flk-1 indicated that the addition of six nucleotides to the 5' end of the isolated KDR clones would provide for a complete coding region.

To achieve this, an EcoRI-BamHI restriction fragment of BTIV200.2 is cloned into the plasmid pBlueScript KS (Strategene, La Jolla, CA). The 5' end of the inserted DNA is blunt ended with Klenow polymerase and Mung Bean nuclease. Next, the synthetic oligonucleotide TCGACGCGCG ATG GAG (SEQ ID NO. 11) is cloned into this vector. The oligonucleotide contains the sequence ATG GAG in frame with the downstream DNA insert. These nucleotides (ATG GAG) encode the amino acids methionine and glutamic acid, the first two amino acids encoded by the KDR gene. The resulting plasmid vector is designated BTIV140. This plasmid is purified on a CsCl gradient.

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The purified plasmid is designated BTIV169. The insert of BTIV169 contains nucleotides 1-2400 (SEQ ID NO. 7) of the KDR gene. A sample of the plasmid pBlueScript KS which contains the clone BTIV169 has been deposited with the American Type Culture Collection and has been assigned ATCC accession number 75200.

Thus, together the clones BTIII081.8, BTIII129.5 and BTIV169 comprise the entire open reading frame of 4,068 nucleotides for the novel KDR gene. As will be discussed below, the KDR gene expresses the novel KDR receptor which binds specifically to the growth factor VEGF.

DNA sequencing of BTIII081.8, BTIII129.5 and BTIV169 (SEQ ID NO. 7) shows that the newly isolated gene is similar to, but distinct from, previously identified type III RTK. The predicted amino acid sequence (SEQ ID NO. 7) contains several structural features which demonstrate that the novel gene is a type III RTK. These structural features are summarized as follows:

1) A hydropathy plot of the predicted amino acid sequence indicates a single membrane spanning region (see Figure 8). This is characteristic of a type III RTK (Figure 7).

2) The putative amino-terminal 762 amino acid portion of the receptor has structural features of extracellular receptor ligand binding domains (1), including regularly spaced cysteines and 18 potential N-linked glycosylation sites (Figure 7).

3) The predicted amino acid sequence of the carboxy-terminal 530 amino acid portion contains an ATP-binding site at lysine 868, 22 amino acids downstream from the consensus ATP recognition sequence Gly-X-Gly-X-X-Gly (26) (Figure 8).

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4) Within the kinase domain there is a 55-60% identical match in amino acid sequence to three other type III receptor tyrosine kinases: ckit proto-oncogene (SEQ ID NO: 8), CSF-1 (SEQ ID NO: 9) and PDGF (SEQ ID NO: 10) (Figure 9).

5) The predicted kinase domain contains a kinase insert domain of approximately 71 amino acids. As indicated in Figure 9, this portion of the amino acid sequence shares little sequence homology with other type III RTK.

The endothelial cell library can be further screened to isolate the 5' untranslated region and genomic clones can be generated so as to isolate the promoter region for the KDR gene.

In addition to the DNA sequence described for the KDR gene (SEQ ID NO. 7), the present invention further comprises DNA sequences which, by virtue of the redundancy of the genetic code, are biologically equivalent to the sequences which encode for the receptor, that is, these other DNA sequences are characterized by nucleotide sequences which differ from those set forth herein, but which encode a receptor having the same amino acid sequences as those encoded by the DNA sequences set forth herein.

In particular, the invention contemplates those DNA sequences which are sufficiently duplicative of the sequence of SEQ ID NO. 7 so as to permit hybridization therewith under standard high stringency Southern hybridization conditions, such as those described in Sambrook et al. (27), as well as the biologically active proteins produced thereby.

This invention also comprises DNA sequences which encode amino acid sequences which differ from those of the novel receptor, but which are the biological equivalent to those described for the

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receptor. Such amino acid sequences may be said to be biologically equivalent to those of the receptor if their sequences differ only by minor deletions from or conservative substitutions to the receptor sequence, such that the tertiary configurations of the sequences are essentially unchanged from those of the receptor.

For example, a codon for the amino acid alanine, a hydrophobic amino acid, may be substituted by a codon encoding another less hydrophobic residue, such as glycine, or a more hydrophobic residue, such as valine, leucine, or isoleucine. Similarly, changes which result in substitution of one negatively charged residue for another, such as aspartic acid for glutamic acid, or one positively charged residue for another, such as lysine for arginine, as well as changes based on similarities of residues in their hydrophobic index, can also be expected to produce a biologically equivalent product. Nucleotide changes which result in alteration of the N-terminal or C-terminal portions of the protein molecule would also not be expected to alter the activity of the protein. It may also be desirable to eliminate one or more of the cysteines present in the sequence, as the presence of cysteines may result in the undesirable formation of multimers when the protein is produced recombinantly, thereby complicating the purification and crystallization processes. Each of the proposed modifications is well within the routine skill in the art, as is determination of retention of biological activity of the encoded products. Therefore, where the terms "KDR gene" or "KDR protein" are used in either the specification or the claims, each will be understood to encompass all such modifications and variations which result in the production of a biologically equivalent protein.

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In addition to the full length gene and protein, the invention encompasses biologically active fragments of each. By "biologically active" is meant a protein fragment which qualitatively retains the receptor activity of the larger KDR protein, or, in the case of a nucleotide sequence, which encodes such a protein fragment. It also refers, for purposes of antibody production, to fragments which are capable of eliciting production of antibodies capable of binding to the receptor protein.

To determine the size of the mRNA transcribed from the kdp gene, Northern blot hybridization experiments are carried out using an EcoRI/BamHI DNA segment (nucleotides 1510-2417, SEQ ID NO. 7) as a hybridization probe. The DNA used for the probe does not contain any portion of the putative kinase domain, and shares little sequence homology to other tyrosine kinases. The Northern blot analysis (Figure 10) shows that a 7 kb band is visualized in cytoplasmic poly(A)+ RNA of ABAE bovine aortic endothelial cells. This transcript differs in size from previously reported transcripts for known type III RTK (7,18).

The isolated cDNA is significant for several reasons. The cDNA encodes a novel type III receptor tyrosine kinase. The homology between the sequence of this cDNA and that of other receptors, as well as structural properties implied by the predicted amino acid sequence confirm the relationship. Receptors for growth factors should have tremendous utility in drug development as they face the outside of the cell and thus are among the best targets for drugs. In addition, the cellular levels of some receptors, in particular the neu proto-oncogene, increase during some cancers. This has been taken advantage of in designing diagnostic tests for these cancers.

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Southern analysis demonstrates that the kdp gene is present in mouse as well as human DNA. Mouse and human (Hela cell) DNA, 15 μ g of each, are digested with 10 units of EcoRI and electrophoresed on a 0.7% agarose gel. The DNA is transferred onto nitrocellulose. The filter is hybridized to a [³²P]CTP-labelled cDNA probe made by nick translating an EcoRI/BamHI fragment from the 5' end of the kdp cDNA (nucleotides 1510-2417, SEQ ID NO. 7). Hybridization is conducted at 30°C in 5 X SSPE, 50% formamide, 0.1% SDS, plus 150 μ g/ml salmon sperm DNA. The DNA probe hybridizes to Southern blots containing EcoRI digested DNA. After 48 hours, the filter is washed at room temperature in 2 X SSC plus 0.1% SDS for 20 minutes, followed by two 20 minute washes at 40°C with 0.1 X SSC plus 0.1% SDS. Autoradiography is then performed for 48 hours. As shown in Figure 11, radioactively labelled DNA is present in both human and mouse samples. This indicates that the kdp gene is present in both species.

An experiment is conducted to ascertain the genetic locus of kdp on human chromosomes. Thirty-eight cell hybrids from 18 unrelated human cell lines and four mouse cell lines are examined (19). A DNA probe hybridizes to Southern blots which contain EcoRI digested DNA from the human-mouse hybrids (using the procedure and DNA probe for human and mouse tissue described in relation to Figure 11). Table I sets forth the results of the segregation of kdp with human chromosomes in EcoRI digested human-mouse somatic cell hybrid DNA:

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Table I

5	<u>Chromosome</u>	Concordant # of Hybrids		Discordant # of Hybrids		<u>% Discordancy</u>
		<u>(+/+)</u>	<u>(-/-)</u>	<u>(+/-)</u>	<u>(-/+)</u>	
	1	4	19	8	4	34
	2	8	18	5	6	30
	3	11	12	3	9	34
10	4	14	24	0	0	0
	5	7	14	7	10	45
	6	7	19	7	5	32
	7	11	14	3	8	31
	8	8	11	6	13	50
15	9	3	20	10	4	38
	10	12	9	2	14	43
	11	9	13	4	11	41
	12	9	10	5	14	50
	13	7	18	7	6	34
20	14	11	8	3	16	50
	15	9	15	5	8	35
	16	7	19	7	5	32
	17	12	7	2	16	49
	18	11	14	3	10	34
25	19	7	18	7	6	34
	20	9	10	5	14	50
	21	11	9	3	15	47
	22	3	16	10	7	47
30	X	8	10	3	8	38

The scoring is determined by the presence(+) or absence (-) of human bands in the hybrids on Southern blots prepared in a similar to those shown in Figure 11. The scoring is compared to the presence or absence of human chromosomes in each hybrid. A 0%

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discordancy indicates a matched segregation of the DNA probe with a chromosome. Three fragments, approximately 6.5 kb, 3.1 kb, and 0.7 kb in size are detected in digests of human DNA (Figure 11), and in all hybrids which had retained human chromosome 4 (Table I). All other chromosomes are excluded in at least 11 discordant hybrids (Table I). The results of Figure 11 and Table I demonstrate that the genetic locus of kdp is on human chromosome 4.

It is noteworthy that both the ckit (3) and the type A PDGF (28) receptor genes map to human chromosome 4. The finding that the genetic locus of kdp is on human chromosome 4 provides further evidence that the novel receptor of this invention is a type III receptor tyrosine kinase.

The next step after identifying the entire coding portion of the kdp gene is to express the receptor protein encoded by that gene. The receptor protein is then utilized so as to identify the growth factor which binds specifically to the receptor.

The receptor protein is expressed using established recombinant DNA methods. Suitable host organisms include bacteria, viruses, yeast, insect or mammalian cell lines, as well as other conventional organisms. For example, CMT-3 monkey kidney cells are transfected with a vector containing the complete coding region of the KDR gene.

The complete coding portion of the KDR gene is assembled by sequentially cloning into pUC119 three DNA fragments derived from BTIII081.8, BTIII129.5, and BTIV169. First, a SmaI-EcoRI fragment of clone BTIII129.5 (nucleotides 3152-4236, SEQ ID NO. 7) is blunt ended with Klenow polymerase and introduced into a SmaI site in pUC119. Next, a BamHI-SmaI fragment of clone BTIII081.8 (nucleotides 2418-3151, SEQ ID NO. 7)

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is introduced at a BamHI-SmaI site. Finally, a SalI-BamHI fragment of clone BTIV169 (nucleotides 1-2417, SEQ ID NO. 7) is introduced at a SalI-BamHI site. Part of the cloning site of pUC119 is contained
5 in the SalI-BamHI fragment, 5' to the KDR gene. In order to clone the complete coding portion into an expression vector, the assembled DNA (in pUC119) is digested with SalI and Asp118 and recloned into the eukaryotic expression vector pcDNAlt_{tkp}ASP.

10 This vector is a modification of the vector pcDNA1 (Invitrogen; San Diego, CA). Specifically, the ampicillin resistance gene is cloned from pBR322 into pcDNA1. A small SV40 T splice and the SV40 polyadenylation signal are then removed and are replaced with a
15 Herpes Simplex Virus-1 polyadenylation signal. Finally, a cytomegalovirus intermediate early splice is inserted 5' to the cloning site to yield pcDNAlt_{tkp}ASP.

Transfection of CMT-3 cells is done using DEAE-dextran. Forty-eight hours after transfection,
20 expression of the novel receptor is monitored using Western blot analysis as follows.

An antibody is used to assay the expressed receptor protein. The predicted amino acid sequence of the receptor is used to generate peptide-derived
25 antibodies to the receptor by conventional techniques. The presence of the novel receptor protein is confirmed by Western blot hybridization.

Specifically, a synthetic peptide with 13 residues is synthesized based on the 12 residues
30 corresponding to amino acids 986-997 of the putative amino acid sequence of the KDR protein (SEQ ID NO. 7), with a cysteine residue linked to the lysine (amino acid 997). The cysteine facilitates coupling of the peptide to a macromolecule which functions as a carrier
35 for the peptide. For example, the peptide is coupled

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to keyhole limpet haemocyanin (KLH) using m-maleimido-benzoyl-N-hydroxysuccinimide ester. Other conventional carriers may be used such as human and bovine serum albumins, myoglobins, β -galactosidase, penicillinase and bacterial toxoids, as well as synthetic molecules such as multi-poly-DL-alanyl-poly-L-lysine and poly-L-lysine.

Rabbits are immunized with the peptide-KLH conjugate to raise polyclonal antibodies. After different periods of time, serum is collected from the rabbits. The IgG fraction of the serum is then purified using a protein A Sepharose column (Pharmacia LKB, Uppsala, Sweden) to obtain the antibody which is designated anti-KDR.PS23.

A sample of the expressed KDR protein is subjected to SDS-PAGE using a 7% acrylamide gel under standard conditions. The protein band is then transferred on to nitrocellulose paper for Western blot analysis and the anti-KDR.PS23 antibody is added at a dilution of 1:1,000 to allow the antibody to react with the protein present. A second antibody, goat anti-rabbit antibody to rabbit IgG, which binds to anti-KDR.PS23, is then added. The detection of proteins which react with the antibodies is performed by autoradiography of bands using an ECL system (Amersham, Chicago, IL). The results are depicted in Figure 12.

Figure 12 shows that a 190 kD protein is present in the cells transfected with the vector containing the KDR gene, but is absent in cells transfected with vector alone. The size of this protein is consistent with it being encoded by the KDR gene, in that the predicted amino acid sequence for the unglycosylated KDR protein is 156 kD, and that sequence contains 18 putative extracellular glycosylation sites

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which would account for the balance of the size seen in the 190 kD band.

5 The expressed receptor is then used to identify the growth factor which interacts with the receptor. In order to test the hypothesis that the KDR protein is a receptor for VEGF, radioligand binding studies are performed. VEGF (provided by D. Gospodarowicz) is radiolabelled with ^{125}I . Cells are
10 transfected with either the vector pcDNA1tkpASP alone (bars 1 and 2 of Figure 13) or with the vector containing the KDR gene (bars 3 and 4). Forty-eight hours later, the transfected cell samples are washed with PBS and then incubated for 90 minutes with serum-free media containing 50 pM [^{125}I]VEGF (specific activity equal to 4,000 cpm per fmol). Excess
15 nonradioactive VEGF, 5 nM, is added to some samples (bars 2 and 4) to define specific binding sites. The samples are washed with ice cold PBS, and the cells are transferred to gamma-counting tubes using a detergent, 0.1% lubrol.
20

The results of the radioligand binding studies are depicted in Figure 13. Figure 13 shows that CMT-3 cells transfected with vector containing the KDR gene contain specific binding sites for [^{125}I]VEGF (compare bars 3 and 4), while cells transfected with
25 vector alone do not (compare bars 1 and 2).

Further evidence that the KDR gene encodes a receptor for VEGF is demonstrated by affinity cross-linking studies (Figure 14). Figure 14 depicts
30 the results of affinity cross-linking of [^{125}I]VEGF to CMT-3 cells which express the KDR protein. CMT-3 cells are transfected with either the pcDNA1tkpASP vector alone (lane 1 of Figure 14) or with the vector containing the KDR gene (lane 2). Forty-eight hours
35 later, the cells are washed in PBS, and serum free

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media containing 200 pM [125 I]VEGF is added. After 90 minutes at room temperature, an affinity cross-linker disuccinimidyl suberate (Pierce Biochemicals, Rockford, IL), 0.5mM, is added for 15 minutes. The samples are then subjected to SDS-PAGE autoradiography.

Three protein bands are seen in SDS-PAGE autoradiograms from samples of CMT-3 cells transfected with the KDR gene and cross-linked to [125 I]VEGF (lane 1). The size of band 1 (235 kD) is consistent with it being the 190 kD protein seen by Western blot analysis (Figure 12), because a 45 kD [125 I] VEGF dimer plus 190 kD would migrate in a manner identical to band 1. The origin of band 2 is not clear, but may represent an altered glycosylation form of band 1. Band 3 (22.5 kD) is most likely VEGF itself, and can be seen faintly in cells transfected with vector alone (lane 2).

The novel KDR gene of this invention is significant for several reasons. Studies of the cellular mechanisms by which receptors function in signal transduction have led in the past to a better understanding of how cells grow in both normal and diseased states. Receptor tyrosine kinases, in particular, have received a great deal of attention because of the observation that a number of RTK are the cellular counterparts for viral oncogenes, implying a direct correlation between changes in the expression of RTK and cancer. In view of this, it is likely that pharmaceuticals targeted at RTK will inhibit the changes in cell growth associated with cancer. In addition, it is likely that monitoring the levels of expression of RTK will prove valuable in diagnosing the onset of cancer.

The described cDNA is isolated from a human endothelial cell library. Endothelial cells participate in angiogenesis, the formation of new blood

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capillaries. Previous work directed towards identifying the growth factors which regulate angiogenesis have primarily focused upon FGF (13), although recent evidence has indicated that other growth factors may be involved as well (12,15,29). This evidence consists of the observations that: 1) FGF does not contain a signal sequence (24) and thus may not be secreted from cells in a manner consistent with the tight regulation of angiogenesis, and 2) endothelial cells synthesize FGF and yet are normally resting (15). Our discovery, then, of a novel growth factor receptor may ultimately clarify these inconsistencies and lead to a better understanding of endothelial cell function.

The teachings of this invention can be readily used by those skilled the art for the purpose of testing pharmaceuticals targeted at the KDR protein. Two examples of approaches which can be used for this purpose are now given.

First, the methods described in this invention for studying the interaction of VEGF with KDR protein can be used to test for pharmaceuticals which will antagonize that interaction. For these studies, cells expressing the KDR protein are incubated with [¹²⁵I]VEGF, together with a candidate pharmaceutical. Inhibition of radioligand binding is tested for; significant inhibition indicates the candidate is an antagonist. Permanent expression of the KDR protein in a cell type such as NIH3T3 cells would make these studies less laborious. This can be easily achieved by those skilled in the art using the described methods.

Second, using the teachings of this invention, those skilled in the art can study structural properties of the KDR protein involved in receptor function. This structural information can

- 30 -

then be used to more rationally design pharmaceuticals
which inhibit that function. Mutagenesis of the KDR
gene by well established protocols is one approach,
crystallization of the receptor binding site is
another.

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- 33 -

SEQUENCE LISTING

(1) GENERAL INFORMATION:

5

(i) APPLICANT: Terman, Bruce I
Carrion, Miguel E

(ii) TITLE OF INVENTION: Identification of a
Novel Human Growth Factor Receptor

10

(iii) NUMBER OF SEQUENCES: 14

(iv) CORRESPONDENCE ADDRESS:

15

(A) ADDRESSEE: Alan M. Gordon
American Cyanamid Company

(B) STREET: 1937 West Main Street, P.O. Box 60

20

(C) CITY: Stamford

(D) STATE: Connecticut

(E) COUNTRY: USA

25

(F) ZIP: 06904

(v) COMPUTER READABLE FORM:

30

(A) MEDIUM TYPE: Floppy disk

(B) COMPUTER: IBM PC AT

(C) OPERATING SYSTEM: MS-DOS

35

- 34 -

(D) SOFTWARE: ASCII from IBM DW 4

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5 (A) APPLICATION NUMBER:

(B) FILING DATE:

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(A) NAME: Gordon, Alan M.

20

(B) REGISTRATION NUMBER: 30,637

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25 (ix) TELECOMMUNICATION INFORMATION:

(A) TELEPHONE: 203 321 2719

(B) TELEFAX: 203 321 2971

30

(C) TELEX:

(2) INFORMATION FOR SEQ ID NO: 1:

35 (i) SEQUENCE CHARACTERISTICS:

- 35 -

(A) LENGTH: 27 base pairs

(B) TYPE: nucleic acid

5

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

10

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

GTCGAC AAY CTG TTG GGR GCC TGC AAC 27

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(2) INFORMATION FOR SEQ ID NO: 2 :

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 35 base pairs

(B) TYPE: nucleic acid

25

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

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(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

GAATTC AG CAC KTT NCT RGC YGC CAG GTC TGY GTC 35

35

- 36 -

(2) INFORMATION FOR SEQ ID NO: 3:

(i) SEQUENCE CHARACTERISTICS:

- 5 (A) LENGTH: 363 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- 10 (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

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AGG ACG AAG AGA AAT GAA TTT GTC CCC TAC AAG ACC 72

20 AAA GGG GCA CGA TTC CGT CAA GGG AAA GAC TAC GTT 108

GGA GCA ATC CCT GTG GAT CTG AAA CGG CGC TTG GAC 144

25 ACG CAT CAC CAG TAG CCA GAG CTC AGC CAG CTC TGG 180

ATT TGT GGA GGA GAA GTC CCT CAG TGA TGT AGA AGA 216

AGA GGA AGC TCC TGA AGA TCT GTA TAA GGA CTT CCT 252

30 GAC CTT GGA GCA TCT CAT CTG TTA CAG TTT CCA AGT 288

GGC TAA GGG CAT GGA GTT CTT GGC ATC GCG AAA GTG 324

35 TAT CCA CAG AGA CCT GGC AGC CAG GAA CGT GCT GAA 360

- 37 -

TTC

363

(2) INFORMATION FOR SEQ ID NO: 4 :

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 251 base pairs

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(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

15

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

20

GTC GAC AAT CTG TTG GGG GCC TGC ACC ATC CCA ACA 36

TCC TGC TGC TCT ACA ACT ATT TTT ATG ACC GGA GGA 72

GGA TCT ACT TGA TTC TAG AGT ATG CCC CCC GCG GAG 108

25

CTC TAC AAG GAG CTG CAG AAG AGC TGC ACA TTT GAC 144

GAG CAG CGA ACA GCC ACG ATC ATG GAG GAG TTG GCA 180

30

GAT GCT CTA ATG TAC TGC CGT GGG AAG AAG GTG ATT 216

CAC AGA GAC CTG GCA GCC AGC AAC GTG CTG AAT TC 251

35

(2) INFORMATION FOR SEQ ID NO: 5:

- 38 -

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 510 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(ix) FEATURE:

(A) NAME/KEY: PDGF Receptor DNA

(B) LOCATION: Internal sequence

(x) PUBLICATION INFORMATION:

(A) AUTHORS: Gronwald, R., et al.

(B) JOURNAL: Proc. Natl. Acad. Sci.

(C) VOLUME: 85

(D) PAGES: 3435-3439

(E) DATE: 1988

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

AAC CTG TGG GGG CCT GCA CCA AAG GAG GAC CAT CTA

36

- 39 -

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30
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TAT CAT CTA TAT CAT CAC TGA GTA CTG CCG CTA CGG 72
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CAG CGC GGA GCT CTA CAG CAA TGC TCT GCC CGT TGG 180
GCT CCC CCT GCC CAG CCA TGT GTC CTT GAC CGG GGG 216
AGA GCG ACG GTG GCT ACA TGG ACA TGA GCA AGG ACG 252
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CAT GGC CCC TTA CGA TAA CTA CGT TCC CTC TGC CCC 360
TGA GAG GAC CTG CCG AGC AAC TTT GAT CAA CGA GTC 396
TCC AGT GCT AAG CTA CAT GGA CCT CGT GGG CTT CAG 432
CTA CCA GGT GGC CAA TGG CAT GGA GTT CTG GCC TCC 468
AAG AAC TGC GTC CAC AGA GAC CTG GCG GCT AGG AAC 504
GTC CTT 510

(2) INFORMATION FOR SEQ ID NO: 6:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 255 base pairs

- 40 -

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

5 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(ix) FEATURE:

10

(A) NAME/KEY: FGF Receptor DNA

(B) LOCATION: Internal sequence

15 (x) PUBLICATION INFORMATION:

(A) AUTHORS: Ruta, M., et al.

(B) JOURNAL: Oncogene

20

(C) VOLUME: 3

(D) PAGES: 9-15

25

(E) DATE: 1988

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

30

AAC CTG CTG GGG GCC TGC ACG CAG GAT GGT CCC TTG 36

TAT GTC ATC GTG GAG TAT GCC TCC AAG GGC AAC CTG 72

CGG GAG TAC CTG CAG ACC CGG AGG CCC CCA GGG CTG 108

35

GAA TAC TGC TAT AAC CCC AGC CAC AAC CCA GAG GAG 144

- 41 -

CAG CTC TCC TCC AAG GAC CTG GTG TCC TGC GCC TAC 180

CAG GAG GCC CGA GGC ATG GAG TAT CTG GCC TCC AAG 216

5 AAG TGC ATA CAC CGA GAC CTG GCA GCC AGG AAT GTC 252

CTG 255

(2) INFORMATION FOR SEQ ID NO: 7:

10

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 4236 base pairs

15

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

20

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

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30 TGG CTC TGC GTG GAG ACC CGG GCC GCC TCT GTG GGT 69

Trp Leu Cys Val Glu Thr Arg Ala Ala Ser Val Gly

15

20

35 TTG CCT AGT GTT TCT CTT GAT CTG CCC AGG CTC AGC 105

Leu Pro Ser Val Ser Leu Asp Leu Pro Arg Leu Ser

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- 42 -

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	40 45	
5	ACT CTT CAA ATT ACT TGC AGG GGA CAG AGG GAC TTG	177
	Thr Leu Gln Ile Thr Cys Arg Gly Gln Arg Asp Leu	
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10	GAC TGG CTT TGG CCC AAT AAT CAG AGT GGC AGT GAG	213
	Asp Trp Leu Trp Pro Asn Asn Gln Ser Gly Ser Glu	
	60 65 70	
15	CAA AGG GTG GAG GTG ACT GAG TGC AGC GAT GGC CTC	249
	Gln Arg Val Glu Val Thr Glu Cys Ser Asp Gly Leu	
	75 80	
20	TTC TGT AAG ACA CTC ACA ATT CCA AAA GTG ATC GGA	285
	Phe Cys Lys Thr Leu Thr Ile Pro Lys Val Ile Gly	
	85 90 95	
25	AAT GAC ACT GGA GCC TAC AAG TGC TTC TAC CGG GAA	321
	Asn Asp Thr Gly Ala Tyr Lys Cys Phe Tyr Arg Glu	
	100 105	
30	ACT GAC TTG GCC TCG GTC ATT TAT GTC TAT GTT CAA	357
	Thr Asp Leu Ala Ser Val Ile Tyr Val Tyr Val Gln	
	110 115	
35	GAT TAC AGA TCT CCA TTT ATT GCT TCT GTT AGT GAC	393
	Asp Tyr Arg Ser Pro Phe Ile Ala Ser Val Ser Asp	
	120 125 130	
40	CAA CAT GGA GTC GTG TAC ATT ACT GAG AAC AAA AAC	429
	Gln His Gly Val Val Tyr Ile Thr Glu Asn Lys Asn	
	135 140	

- 43 -

	AAA ACT GTG GTG ATT CCA TGT CTC GGG TCC ATT TCA	465
	Lys Thr Val Val Ile Pro Cys Leu Gly Ser Ile Ser	
	145 150 155	
5	AAT CTC AAC GTG TCA CTT TGT GCA AGA TAC CCA GAA	501
	Asn Leu Asn Val Ser Leu Cys Ala Arg Tyr Pro Glu	
	160 165	
10	AAG AGA TTT GTT CCT GAT GGT AAC AGA ATT TCC TGG	537
	Lys Arg Phe Val Pro Asp Gly Asn Arg Ile Ser Trp	
	170 175	
15	GAC AGC AAG AAG GGC TTT ACT ATT CCC AGC TAC ATG	573
	Asp Ser Lys Lys Gly Phe Thr Ile Pro Ser Tyr Met	
	180 185 190	
20	ATC AGC TAT GCT GGC ATG GTC TTC TGT GAA GCA AAA	609
	Ile Ser Tyr Ala Gly Met Val Phe Cys Glu Ala Lys	
	195 200	
25	ATT AAT GAT GAA AGT TAC CAG TCT ATT ATG TAC ATA	645
	Ile Asn Asp Glu Ser Tyr Gln Ser Ile Met Tyr Ile	
	205 210 215	
30	GTT GTC GTT GTA GGG TAT AGG ATT TAT GAT GTG GTT	681
	Val Val Val Val Gly Tyr Arg Ile Tyr Asp Val Val	
	220 225	
35	CTG AGT CCG TCT CAT GGA ATT GAA CTA TCT GTT GGA	717
	Leu Ser Pro Ser His Gly Ile Glu Leu Ser Val Gly	
	230 235	
40	GAA AAG CTT GTC TTA AAT TGT ACA GCA AGA ACT GAA	753
	Glu Lys Leu Val Leu Asn Cys Thr Ala Arg Thr Glu	
	240 245 250	

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	CTA AAT GTG GGG ATT GAC TTC AAC TGG GAA TAC CCT	789
	Leu Asn Val Gly Ile Asp Phe Asn Trp Glu Tyr Pro	
	255 260	
5	TCT TCG AAG CAT CAG CAT AAG AAA CTT GTA AAC CGA	825
	Ser Ser Lys His Gln His Lys Lys Leu Val Asn Arg	
	265 270 275	
10	GAC CTA AAA ACC CAG TCT GGG AGT GAG ATG AAG AAA	861
	Asp Leu Lys Thr Gln Ser Gly Ser Glu Met Lys Lys	
	280 285	
15	TTT TTG AGC ACC TTA ACT ATA GAT GGT GTA ACC CGG	897
	Phe Leu Ser Thr Leu Thr Ile Asp Gly Val Thr Arg	
	290 295	
20	AGT GAC CAA GGA TTG TAC ACC TGT GCA GCA TCC AGT	933
	Ser Asp Gln Gly Leu Tyr Thr Cys Ala Ala Ser Ser	
	300 305 310	
	GGG CTG ATG ACC AAG AAG AAC AGC ACA TTT GTC AGG	969
	Gly Leu Met Thr Lys Lys Asn Ser Thr Phe Val Arg	
	315 320	
25	GTC CAT GAA AAA CCT TTT GTT GCT TTT GGA AGT GGC	1005
	Val His Glu Lys Pro Phe Val Ala Phe Gly Ser Gly	
	325 330 335	
30	ATG GAA TCT CTG GTG GAA GCC ACG GTG GGG GAG CGT	1041
	Met Glu Ser Leu Val Glu Ala Thr Val Gly Glu Arg	
	340 345	
35	GTC AGA ATC CCT GCG AAG TAC CTT GGT TAC CCA CCC	1077
	Val Arg Ile Pro Ala Lys Tyr Leu Gly Tyr Pro Pro	
	350 355	

- 45 -

	CCA GAA ATA AAA TGG TAT AAA AAT GGA ATA CCC CTT	1113
	Pro Glu Ile Lys Trp Tyr Lys Asn Gly Ile Pro Leu	
	360 365 370	
5	GAG TCC AAT CAC ACA ATT AAA GCG GGG CAT GTA CTG	1149
	Glu Ser Asn His Thr Ile Lys Ala Gly His Val Leu	
	375 380	
10	ACG ATT ATG GAA GTG AGT GAA AGA GAC ACA GGA AAT	1185
	Thr Ile Met Glu Val Ser Glu Arg Asp Thr Gly Asn	
	385 390 395	
15	TAC ACT GTC ATC CTT ACC AAT CCC ATT TCA AAG GAG	1221
	Tyr Thr Val Ile Leu Thr Asn Pro Ile Ser Lys Glu	
	400 405	
20	AAG CAG AGC CAT GTG GTC TCT CTG GTT GTG TAT GTC	1257
	Lys Gln Ser His Val Val Ser Leu Val Val Tyr Val	
	410 415	
25	CCA CCC CAG ATT GGT GAG AAA TCT CTA ATC TCT CCT	1293
	Pro Pro Gln Ile Gly Glu Lys Ser Leu Ile Ser Pro	
	420 425 430	
30	GTG GAT TCC TAC CAG TAC GGC ACC ACT CAA ACG CTG	1329
	Val Asp Ser Tyr Gln Tyr Gly Thr Thr Gln Thr Leu	
	435 440	
35	ACA TGT ACG GTC TAT GCC ATT CCT CCC CCG CAT CAC	1365
	Thr Cys Thr Val Tyr Ala Ile Pro Pro Pro His His	
	445 450 455	
40	ATC CAC TGG TAT TGG CAG TTG GAG GAA GAG TGC GCC	1401
	Ile His Trp Tyr Trp Gln Leu Glu Glu Glu Cys Ala	
	460 465	

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AAC GAG CCC AGC CAA GCT GTC TCA GTG ACA AAC CCA 1437
Asn Glu Pro Ser Gln Ala Val Ser Val Thr Asn Pro
470 475

5 TAC CCT TGT GAA GAA TGG AGA AGT GTG GAG GAC TTC 1473
Tyr Pro Cys Glu Glu Trp Arg Ser Val Glu Asp Phe
480 485 490

10 CAG GGA GGA AAT AAA ATT GAA GTT AAT AAA AAT CAA 1509
Gln Gly Gly Asn Lys Ile Glu Val Asn Lys Asn Gln
495 500

15 TTT GCT CTA ATT GAA GGA AAA AAC AAA ACT GTA AGT 1545
Phe Ala Leu Ile Glu Gly Lys Asn Lys Thr Val Ser
505 510 515

20 ACC CTT GTT ATC CAA GCG GCA AAT GTG TCA GCT TTG 1581
Thr Leu Val Ile Gln Ala Ala Asn Val Ser Ala Leu
520 525

TAC AAA TGT GAA GCG GTC AAC AAA GTC GGG AGA GGA 1617
Tyr Lys Cys Glu Ala Val Asn Lys Val Gly Arg Gly
530 535

25 GAG AGG GTG ATC TCC TTC CAC GTG ACC AGG GGT CCT 1653
Glu Arg Val Ile Ser Phe His Val Thr Arg Gly Pro
540 545 550

30 GAA ATT ACT TTG CAA CCT GAC ATG CAG CCC ACT GAG 1689
Glu Ile Thr Leu Gln Pro Asp Met Gln Pro Thr Glu
555 560

35 CAG GAG AGC GTG TCT TTG TGG TGC ACT GCA GAC AGA 1725
Gln Glu Ser Val Ser Leu Trp Cys Thr Ala Asp Arg
565 570 575

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	TCT ACG TTT GAG AAC CTC ACA TGG TAC AAG CTT GGC	1761
	Ser Thr Phe Glu Asn Leu Thr Trp Tyr Lys Leu Gly	
	580 585	
5	CCA CAG CCT CTG CCA ATC CAT GTG GGA GAG TTG CCC	1797
	Pro Gln Pro Leu Pro Ile His Val Gly Glu Leu Pro	
	590 595	
10	ACA CCT GTT TGC AAG AAC TTG GAT ACT CTT TGG AAA	1833
	Thr Pro Val Cys Lys Asn Leu Asp Thr Leu Trp Lys	
	600 605 610	
15	TTG AAT GCC ACC ATG TTC TCT AAT AGC ACA AAT GAC	1869
	Leu Asn Ala Thr Met Phe Ser Asn Ser Thr Asn Asp	
	615 620	
20	ATT TTG ATC ATG GAG CTT AAG AAT GCA TCC TTG CAG	1905
	Ile Leu Ile Met Glu Leu Lys Asn Ala Ser Leu Gln	
	625 630 635	
	GAC CAA GGA GAC TAT GTC TGC CTT GCT CAA GAC AGG	1941
	Asp Gln Gly Asp Tyr Val Cys Leu Ala Gln Asp Arg	
	640 645	
25	AAG ACC AAG AAA AGA CAT TGC GTG GTC AGG CAG CTC	1977
	Lys Thr Lys Lys Arg His Cys Val Val Arg Gln Leu	
	650 655	
30	ACA GTC CTA GAG CGT GTG GCA CCC ACG ATC ACA GGA	2013
	Thr Val Leu Glu Arg Val Ala Pro Thr Ile Thr Gly	
	660 665 670	
35	AAC CTG GAG AAT CAG ACG ACA AGT ATT GGG GAA AGC	2049
	Asn Leu Glu Asn Gln Thr Thr Ser Ile Gly Glu Ser	
	675 680	

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	ATC GAA GTC TCA TGC ACG GCA TCT GGG AAT CCC CCT	2085
	Ile Glu Val Ser Cys Thr Ala Ser Gly Asn Pro Pro	
	685 690 695	
5	CCA CAG ATC ATG TGG TTT AAA GAT AAT GAG ACC CTT	2121
	Pro Gln Ile Met Trp Phe Lys Asp Asn Glu Thr Leu	
	700 705	
10	GTA GAA GAC TCA GGC ATT GTA TTG AAG GAT GGG AAC	2157
	Val Glu Asp Ser Gly Ile Val Leu Lys Asp Gly Asn	
	710 715	
15	CGG AAC CTC ACT ATC CGC AGA GTG AGG AAG GAG GAC	2193
	Arg Asn Leu Thr Ile Arg Arg Val Arg Lys Glu Asp	
	720 725 730	
20	GAA GGC CTC TAC ACC TGC CAG GCA TGC AGT GTT CTT	2229
	Glu Gly Leu Tyr Thr Cys Gln Ala Cys Ser Val Leu	
	735 740	
25	GGC TGT GCA AAA GTG GAG GCA TTT TTC ATA ATA GAA	2265
	Gly Cys Ala Lys Val Glu Ala Phe Phe Ile Ile Glu	
	745 750 755	
30	GGT GCC CAG GAA AAG ACG AAC TTG GAA ATC ATT ATT	2301
	Gly Ala Gln Glu Lys Thr Asn Leu Glu Ile Ile Ile	
	760 765	
35	CTA GTA GGC ACG ACG GTG ATT GCC ATG TTC TTC TGG	2337
	Leu Val Gly Thr Thr Val Ile Ala Met Phe Phe Trp	
	770 775	
40	CTA CTT CTT GTC ATC ATC CTA GGG ACC GTT AAG CGG	2373
	Leu Leu Leu Val Ile Ile Leu Gly Thr Val Lys Arg	
	780 785 790	

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	GCC AAT GGA GGG GAA CTG AAG ACA GGC TAC TTG TCC	2409
	Ala Asn Gly Gly Glu Leu Lys Thr Gly Tyr Leu Ser	
	795 800	
5	ATC GTC ATG GAT CCA GAT GAA CTC CCA TTG GAT GAA	2445
	Ile Val Met Asp Pro Asp Glu Leu Pro Leu Asp Glu	
	805 810 815	
10	CAT TGT GAA CGA CTG CCT TAT GAT GCC AGC AAA TGG	2481
	His Cys Glu Arg Leu Pro Tyr Asp Ala Ser Lys Trp	
	820 825	
15	GAA TTC CCC AGA GAC CGG CTG AAC CTA GGT AAG CCT	2517
	Glu Phe Pro Arg Asp Arg Leu Asn Leu Gly Lys Pro	
	830 835	
20	CTT GGC CGT GGT GCC TTT GGC CAA GAG ATT GAA GCA	2553
	Leu Gly Arg Gly Ala Phe Gly Gln Glu Ile Glu Ala	
	840 845 850	
	GAT GCC TTT GGA ATT GAC AAG ACA GCA ACT TGC AGG	2589
	Asp Ala Phe Gly Ile Asp Lys Thr Ala Thr Cys Arg	
	855 860	
25	ACA GTA GCA GTC AAA ATG TTG AAA GAA GGA GCA ACA	2625
	Thr Val Ala Val Lys Met Leu Lys Glu Gly Ala Thr	
	865 870 875	
30	CAC AGT GAG CAT CGA GCT CTC ATG TCT GAA CTC AAG	2661
	His Ser Glu His Arg Ala Leu Met Ser Glu Leu Lys	
	880 885	
35	ATC CTC ATT CAT ATT GGT CAC CAT CTC AAT GTG GTC	2697
	Ile Leu Ile His Ile Gly His His Leu Asn Val Val	
	890 895	

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	AAC CTT CTA GGT GCC TGT ACC AAG CCA GGA GGG CCA	2733
	Asn Leu Leu Gly Ala Cys Thr Lys Pro Gly Gly Pro	
	900 905 910	
5	CTC ATG GTG ATT GTG GAA TTC TGC AAA TTT GGA AAC	2769
	Leu Met Val Ile Val Glu Phe Cys Lys Phe Gly Asn	
	915 920	
10	CTG TCC ACT TAC CTG AGG AGC AAG AGA AAT GAA TTT	2805
	Leu Ser Thr Tyr Leu Arg Ser Lys Arg Asn Glu Phe	
	925 930 935	
15	GTC CCC TAC AAG ACC AAA GGG GCA CGA TTC CGT CAA	2841
	Val Pro Tyr Lys Thr Lys Gly Ala Arg Phe Arg Gln	
	940 945	
20	GGG AAA GAC TAC GTT GGA GCA ATC CCT GTG GAT CTG	2877
	Gly Lys Asp Tyr Val Gly Ala Ile Pro Val Asp Leu	
	950 955	
25	AAA CGG CGC TTG GAC AGC ATC ACC AGT AGC CAG AGC	2913
	Lys Arg Arg Leu Asp Ser Ile Thr Ser Ser Gln Ser	
	960 965 970	
30	TCA GCC AGC TCT GGA TTT GTG GAG GAG AAG TCC CTC	2949
	Ser Ala Ser Ser Gly Phe Val Glu Glu Lys Ser Leu	
	975 980	
35	AGT GAT GTA GAA GAA GAG GAA GCT CCT GAA GAT CTG	2985
	Ser Asp Val Glu Glu Glu Glu Ala Pro Glu Asp Leu	
	985 990 995	
40	TAT AAG GAC TTC CTG ACC TTG GAG CAT CTC ATC TGT	3021
	Tyr Lys Asp Phe Leu Thr Leu Glu His Leu Ile Cys	
	1000 1005	

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TAC AGC TTC CAA GTG GCT AAG GGC ATG GAG TTC TTG 3057
 Tyr Ser Phe Gln Val Ala Lys Gly Met Glu Phe Leu
 1010 1015

5 GCA TCG CGA AAG TGT ATC CAC AGG GAC CTG GCG GCA 3093
 Ala Ser Arg Lys Cys Ile His Arg Asp Leu Ala Ala
 1020 1025 1030

10 CGA AAT ATC CTC TTA TCG GAG AAG AAC GTG GTT AAA 3129
 Arg Asn Ile Leu Leu Ser Glu Lys Asn Val Val Lys
 1035 1040

15 ATC TGT GAC TTT GGC TTG GCC CGG GAT ATT TAT AAA 3165
 Ile Cys Asp Phe Gly Leu Ala Arg Asp Ile Tyr Lys
 1045 1050 1055

20 GAT CCA GAT TAT GTC AGA AAA GGA GAT GCT CGC CTC 3201
 Asp Pro Asp Tyr Val Arg Lys Gly Asp Ala Arg Leu
 1060 1065

CCT TTG AAA TGG ATG GCC CCA GAA ACA ATT TTT GAC 3237
 Pro Leu Lys Trp Met Ala Pro Glu Thr Ile Phe Asp
 1070 1075

25 AGA GTG TAC ACA ATC CAG AGT GAC GTC TGG TCT TTT 3273
 Arg Val Tyr Thr Ile Gln Ser Asp Val Trp Ser Phe
 1080 1085 1090

30 GGT GTT TTG CTG TGG GAA ATA TTT TCC TTA GGT GCT 3309
 Gly Val Leu Leu Trp Glu Ile Phe Ser Leu Gly Ala
 1095 1100

35 TCT CCA TAT CCT GGG GTA AAG ATT GAT GAA GAA TTT 3345
 Ser Pro Tyr Pro Gly Val Lys Ile Asp Glu Glu Phe
 1105 1110 1115

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	TGT AGG CGA TTG AAA GAA GGA ACT AGA ATG AGG GCC	3381
	Cys Arg Arg Leu Lys Glu Gly Thr Arg Met Arg Ala	
	1120 1125	
5	CCT GAT TAT ACT ACA CCA GAA ATG TAC CAG ACC ATG	3417
	Pro Asp Tyr Thr Thr Pro Glu Met Tyr Gln Thr Met	
	1130 1135	
10	CTG GAC TGC TGG CAC GGG GAG CCC AGT CAG AGA CCC	3453
	Leu Asp Cys Trp His Gly Glu Pro Ser Gln Arg Pro	
	1140 1145 1150	
15	ACG TTT TCA GAG TTG GTG GAA CAT TTG GGA AAT CTC	3489
	Thr Phe Ser Glu Leu Val Glu His Leu Gly Asn Leu	
	1155 1160	
20	TTG CAA GCT AAT GCT CAG CAG GAT GGC AAA GAC TAC	3525
	Leu Gln Ala Asn Ala Gln Gln Asp Gly Lys Asp Tyr	
	1165 1170 1175	
25	ATT GTT CTT CCG ATA TCA GAG ACT TTG AGC ATG GAA	3561
	Ile Val Leu Pro Ile Ser Glu Thr Leu Ser Met Glu	
	1180 1185	
30	GAG GAT TCT GGA CTC TCT CTG CCT ACC TCA CCT GTT	3597
	Glu Asp Ser Gly Leu Ser Leu Pro Thr Ser Pro Val	
	1190 1195	
35	TCC TGT ATG GAG GAG GAG GAA GTA TGT GAC CCC AAA	3633
	Ser Cys Met Glu Glu Glu Glu Val Cys Asp Pro Lys	
	1200 1205 1210	
35	TTC CAT TAT GAC AAC ACA GCA GGA ATC AGT CAG TAT	3669
	Phe His Tyr Asp Asn Thr Ala Gly Ile Ser Gln Tyr	
	1215 1220	

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CTG CAG AAC AGT AAG CGA AAG AGC CGG CCT GTG AGT 3705
 Leu Gln Asn Ser Lys Arg Lys Ser Arg Pro Val Ser
 1225 1230 1235

5 GTA AAA ACA TTT GAA GAT ATC CCG TTA GAA GAA CCA 3741
 Val Lys Thr Phe Glu Asp Ile Pro Leu Glu Glu Pro
 1240 1245

10 GAA GTA AAA GTA ATC CCA GAT GAC AAC CAG ACG GAC 3777
 Glu Val Lys Val Ile Pro Asp Asp Asn Gln Thr Asp
 1250 1255

15 AGT GGT ATG GTT CTT GCC TCA GAA GAG CTG AAA ACT 3813
 Ser Gly Met Val Leu Ala Ser Glu Glu Leu Lys Thr
 1260 1265 1270

20 TTG GAA GAC AGA ACC AAA TTA TCT CCA TCT TTT GGT 3849
 Leu Glu Asp Arg Thr Lys Leu Ser Pro Ser Phe Gly
 1275 1280

GGA ATG GTG CCC AGC AAA AGC AGG GAG TCT GTG GCA 3885
 Gly Met Val Pro Ser Lys Ser Arg Glu Ser Val Ala
 1285 1290 1295

25 TCT GAA GGC TCA AAC CAG ACA AGC GGC TAC CAG TCC 3921
 Ser Glu Gly Ser Asn Gln Thr Ser Gly Tyr Gln Ser
 1300 1305

30 GGA TAT CAC TCC GAT GAC ACA GAC ACC ACC GTG TAC 3957
 Gly Tyr His Ser Asp Asp Thr Asp Thr Thr Val Tyr
 1310 1315

35 TCC AGT GAG GAA GCA GAA CTT TTA AAG CTG ATA GAG 3993
 Ser Ser Glu Glu Ala Glu Leu Leu Lys Leu Ile Glu
 1320 1325 1330

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ATT GGA GTG CAA ACC GGT AGC ACA GCC CAG ATT CTC 4029
Ile Gly Val Gln Thr Gly Ser Thr Ala Gln Ile Leu
1335 1340

5 CAG CCT GAC ACG GGG ACC ACA CTG AGC TCT CCT CCT 4065
Gln Pro Asp Thr Gly Thr Thr Leu Ser Ser Pro Pro
1345 1350 1355

10 GTT TAAAAGGAAG CATCCACACC CCAACTCCCG GACATCACAT 4108
Val
1356

GAGAGGTCTG CTCAGATTTT GAAGTGTGT TCTTTCCACC 4148

15 AGCAGGAAGT AGCCGCATTT GATTTTCATT TCGACAACAG 4188

AAAAAGGACC TCGGACTGCA GGGAGCCAGC TCTTCTAGGC 4228

20 TTGTGACC 4236

(2) INFORMATION FOR SEQ ID NO: 8:

(i) SEQUENCE CHARACTERISTICS:

25 (A) LENGTH: 433 amino acids

(B) TYPE: amino acid

(C) STRANDEDNESS:

30

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

35

(ix) FEATURE:

- 55 -

(A) NAME/KEY: ckit proto-oncogene receptor

(B) LOCATION: Amino acids 543-975

5

(x) PUBLICATION INFORMATION:

(A) AUTHORS: Yarden, Y., et al.

10

(B) JOURNAL: EMBO J.

(C) VOLUME: 6

(D) PAGES: 3341-3351

15

(E) DATE: 1987

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

20

Leu Thr Tyr Lys Tyr Leu Gln Lys Pro Met Tyr Glu Val Gln
543 545 550 555

Trp Lys Val Val Glu Glu Ile Asn Gly Asn Asn Tyr Val Tyr
560 565 570

25

Ile Asp Pro Thr Gln Leu Pro Tyr Asp His Lys Trp Glu Phe
575 580

30

Pro Arg Asn Arg Leu Ser Phe Gly Lys Thr Leu Gly Ala Gly
585 590 595

Ala Phe Gly Lys Val Val Ala Glu Thr Ala Tyr Gly Leu Ile
600 605 610

35

Lys Ser Asp Ala Ala Met Thr Val Ala Val Lys Met Leu Lys

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	615		620		625
	Pro Ser Ala His Leu Thr Glu Arg Glu Ala Leu Met Ser Glu				
	630		635		640
5	Leu Lys Val Leu Ser Tyr Leu Gly Asn His Met Asn Ile Val				
	645		650		
	Asn Leu Leu Gly Ala Cys Thr Ile Gly Gly Pro Thr Leu Val				
10	655		660		665
	Ile Thr Glu Tyr Cys Cys Tyr Gly Asp Leu Leu Asn Phe Leu				
	670		675		680
15	Arg Arg Lys Arg Asp Ser Phe Ile Cys Ser Lys Gln Glu Asp				
	685		690		695
	His Ala Glu Ala Ala Leu Tyr Lys Asn Leu Leu His Ser Lys				
	700		705		710
20	Glu Ser Ser Cys Ser Asp Ser Thr Asn Glu Tyr Met Asp Met				
	715		720		
	Lys Pro Gly Val Ser Tyr Val Val Pro Thr Lys Ala Asp Lys				
25	725		730		735
	Arg Arg Ser Val Arg Ile Gly Ser Tyr Ile Glu Arg Asp Val				
	740		745		750
30	Thr Pro Ala Ile Met Glu Asp Asp Glu Leu Ala Leu Asp Leu				
	755		760		765
	Glu Asp Leu Leu Ser Phe Ser Tyr Gln Val Lys Gly Met Ala				
	770		775		780
35					

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Phe Leu Ala Ser Lys Asn Cys Ile His Arg Asp Leu Ala Ala
785 790

5 Arg Asn Ile Leu Leu Thr His Gly Arg Ile Thr Lys Ile Cys
795 800 805

Asp Phe Gly Leu Ala Arg Asp Ile Lys Asn Asp Ser Asn Tyr
810 815 820

10 Val Val Lys Gly Asn Ala Arg Leu Pro Val Lys Val Met Ala
825 830 835

Pro Glu Ser Ile Phe Asn Cys Val Tyr Thr Glu Glu Ser Asp
840 845 850

15 Val Trp Ser Tyr Gly Ile Phe Leu Trp Glu Leu Phe Ser Leu
855 860

Gly Ser Ser Pro Tyr Pro Gly Met Pro Val Lys Ser Lys Phe
20 865 870 875

Tyr Lys Met Ile Lys Glu Gly Phe Arg Met Leu Ser Pro Glu
880 885 890

25 His Ala Pro Ala Glu Met Tyr Asp Ile Met Lys Thr Cys Trp
895 900 905

Asp Ala Asp Pro Leu Lys Arg Pro Thr Phe Lys Gln Ile Val
910 915 920

30 Gln Leu Ile Glu Lys Gln Ile Ser Glu Ser Thr Asn His Ile
925 930

Tyr Ser Asn Leu Ala Asn Cys Ser Pro Asn Arg Gln Lys Pro
35 935 940 945

- 58 -

Val Val Asp His Ser Val Arg Ile Asn Ser Val Gly Ser Thr
950 955 960

Ala Ser Ser Ser Gln Pro Leu Leu Val His Asp Asp Val
965 970 975

5

(2) INFORMATION FOR SEQ ID NO: 9:

(i) SEQUENCE CHARACTERISTICS:

10

(A) LENGTH: 437 amino acids

(B) TYPE: amino acid

15

(C) STRANDEDNESS:

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

20

(ix) FEATURE:

(A) NAME/KEY: CSF-1 receptor

25

(B) LOCATION: Amino acids 536-972

(x) PUBLICATION INFORMATION:

(A) AUTHORS: Coussens, L., et al.

30

(B) JOURNAL: Nature

(C) VOLUME: 320

35

(D) PAGES: 277-280

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(E) DATE: 1986

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

Leu Leu Tyr Lys Tyr Lys Gln Lys Pro Lys Tyr Gln Val Arg
536 540 545

10 Trp Lys Ile Ile Glu Ser Tyr Glu Gly Asn Ser Tyr Thr Phe
550 555 560

Ile Asp Pro Thr Gln Leu Pro Tyr Asn Glu Lys Trp Glu Phe
565 570 575

15 Pro Arg Asn Asn Leu Gln Phe Gly Lys Thr Leu Gly Ala Gly
580 585 590

20 Ala Phe Gly Lys Val Val Glu Ala Thr Ala Phe Gly Leu Gly
595 600 605

Lys Glu Asp Ala Val Leu Lys Val Ala Val Lys Met Leu Lys
610 615

25 Ser Thr Ala His Ala Asp Glu Lys Glu Ala Leu Met Ser Glu
620 625 630

Leu Lys Ile Met Ser His Leu Gly Gln His Glu Asn Ile Val
635 640 645

30 Asn Leu Leu Gly Ala Cys Thr His Gly Gly Pro Val Leu Val
650 655 660

35 Ile Thr Glu Tyr Cys Cys Tyr Gly Asp Leu Leu Asn Phe Leu
665 670 675

- 60 -

Arg Arg Lys Ala Glu Ala Met Leu Gly Pro Ser Leu Ser Pro
680 685

5 Gly Gln Asp Pro Glu Gly Gly Val Asp Tyr Lys Asn Ile His
690 695 700

Leu Glu Lys Lys Tyr Val Arg Arg Asp Ser Gly Phe Ser Ser
705 710 715

10 Gln Gly Val Asp Thr Tyr Val Glu Met Arg Pro Val Ser Thr
720 725 730

Ser Ser Asn Asp Ser Phe Ser Glu Gln Asp Leu Asp Lys Glu
735 740 745

15 Asp Gly Arg Pro Leu Glu Leu Arg Asp Leu Leu His Phe Ser
750 755

Ser Gln Val Ala Gln Gly Met Ala Phe Leu Ala Ser Lys Asn
20 760 765 770

Cys Ile His Arg Asp Val Ala Ala Arg Asn Val Leu Leu Thr
775 780 785

25 Asn Gly His Val Ala Lys Ile Gly Asp Phe Gly Leu Ala Arg
790 795 800

Asp Ile Met Asn Asp Ser Asn Tyr Ile Val Lys Gly Asn Ala
805 810 815

30 Arg Leu Pro Val Lys Trp Met Ala Pro Glu Ser Ile Phe Asp
820 825

Cys Val Tyr Thr Val Gln Ser Asp Val Trp Ser Tyr Gly Ile
35 830 835 840

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Leu Leu Trp Glu Ile Phe Ser Leu Gly Leu Asn Pro Tyr Pro
845 850 855

5 Gly Ile Leu Val Asn Ser Lys Phe Tyr Lys Leu Val Lys Asp
860 865 870

Gly Tyr Gln Met Ala Gln Pro Ala Phe Ala Pro Lys Asn Ile
875 880 885

10 Tyr Ser Ile Met Gln Ala Cys Trp Ala Leu Glu Pro Thr His
890 895

Arg Pro Thr Phe Gln Gln Ile Cys Ser Phe Leu Gln Glu Gln
900 905 910

15 Ala Gln Glu Asp Arg Arg Glu Arg Asp Tyr Thr Asn Leu Pro
915 920 925

Ser Ser Ser Arg Ser Gly Gly Ser Gly Ser Ser Ser Ser Glu
20 930 935 940

Leu Glu Glu Glu Ser Ser Ser Glu His Leu Thr Cys Cys Glu
945 950 955

25 Gln Gly Asp Ile Ala Gln Pro Leu Leu Gln Pro Asn Asn Tyr
960 965

Gln Phe Cys
970

30

(2) INFORMATION FOR SEQ ID NO: 10:

(1) SEQUENCE CHARACTERISTICS:

35 (A) LENGTH: 566 amino acids

- 62 -

(B) TYPE: amino acid

(C) STRANDEDNESS:

5

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

10

(ix) FEATURE:

(A) NAME/KEY: PDGF receptor

(B) LOCATION: Amino acids 522-1087

15

(x) PUBLICATION INFORMATION:

(A) AUTHORS: Gronwald, R., et al.

20

(B) JOURNAL: Proc. Natl. Acad. Sci.

(C) VOLUME: 85

(D) PAGES: 3435-3439

25

(E) DATE: 1988

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:

30

Met	Leu	Trp	Gln	Lys	Lys	Pro	Arg	Tyr	Glu	Ile	Arg	Trp	Lys
522				525				530					535

Val	Ile	Glu	Ser	Val	Ser	Ser	Asp	Gly	His	Glu	Tyr	Ile	Tyr
				540						545			

35

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Val Asp Pro Val Gln Leu Pro Tyr Asp Ser Thr Trp Glu Leu
550 555 560

5 Pro Arg Asp Gln Leu Val Leu Gly Arg Thr Leu Gly Ser Gly
565 570 575

Ala Phe Gly Gln Val Val Glu Ala Thr Ala His Gly Leu Ser
580 585 590

10 His Ser Gln Ala Thr Met Lys Val Ala Val Lys Met Leu Lys
595 600 605

Ser Thr Ala Arg Ser Ser Glu Lys Gln Ser Leu Met Ser Glu
610 615

15 Leu Lys Ile Met Ser His Leu Gly Pro His Leu Asn Val Val
620 625 630

20 Asn Leu Leu Gly Ala Cys Thr Lys Gly Gly Pro Ile Tyr Ile
635 640 645

Ile Thr Glu Tyr Cys Arg Tyr Gly Asp Leu Val Asp Tyr Leu
650 655 660

25 His Arg Asn Lys His Thr Phe Leu Gln Arg His Ser Asn Lys
665 670 675

His Cys Pro Pro Ser Ala Glu Leu Tyr Ser Asn Ala Leu Pro
680 685

30 Val Gly Phe Ser Leu Pro Ser His Leu Asn Leu Thr Gly Glu
690 695 700

35 Ser Asp Gly Gly Tyr Met Asp Met Ser Lys Asp Glu Ser Ile
705 710 715

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Asp Tyr Val Pro Met Leu Asp Met Lys Gly Asp Ile Lys Tyr
720 725 730

5 Ala Asp Ile Glu Ser Pro Ser Tyr Met Ala Pro Tyr Asp Asn
735 740 745

Tyr Val Pro Ser Ala Pro Glu Arg Thr Tyr Arg Ala Thr Leu
750 755

10 Ile Asn Asp Ser Pro Val Leu Ser Tyr Thr Asp Leu Val Gly
760 765 770

Phe Ser Tyr Gln Val Ala Asn Gly Met Asp Phe Leu Ala Ser
775 780 785

15 Lys Asn Cys Val His Arg Asp Leu Ala Ala Arg Asn Val Leu
790 795 800

20 Ile Cys Glu Gly Lys Leu Val Lys Ile Cys Asp Phe Gly Phe
805 810 815

Ala Arg Asp Ile Met Arg Asp Ser Asn Tyr Ile Ser Lys Gly
820 825

25 Ser Thr Tyr Leu Pro Leu Lys Trp Met Ala Pro Glu Ser Ile
830 835 840

Phe Asn Ser Leu Tyr Thr Thr Leu Ser Asp Val Trp Ser Phe
845 850 855

30 Gly Ile Leu Leu Trp Glu Ile Phe Thr Leu Gly Gly Thr Pro
860 865 870

35 Tyr Pro Glu Leu Pro Met Asn Asp Gln Phe Tyr Asn Ala Ile
875 880 885

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Lys Arg Gly Tyr Arg Met Ala Gln Pro Ala His Ala Ser Asp
890 895

5 Glu Ile Tyr Glu Ile Met Gln Lys Cys Trp Glu Glu Lys Phe
900 905 910

Glu Thr Arg Pro Pro Phe Ser Gln Leu Val Leu Leu Leu Glu
915 920 925

10 Arg Leu Leu Gly Glu Gly Tyr Lys Lys Lys Tyr Gln Gln Val
930 935 940

15 Asp Glu Glu Phe Leu Arg Ser Asp His Pro Ala Ile Leu Arg
945 950 955

Ser Gln Ala Arg Phe Pro Gly Ile His Ser Leu Arg Ser Pro
960 965

20 Leu Asp Thr Ser Ser Val Leu Tyr Thr Ala Val Gln Pro Asn
970 975 980

Glu Ser Asp Asn Asp Tyr Ile Ile Pro Leu Pro Asp Pro Lys
985 990 995

25 Pro Asp Val Ala Asp Glu Gly Leu Pro Glu Gly Ser Pro Ser
1000 1005 1010

30 Leu Ala Ser Ser Thr Leu Asn Glu Val Asn Thr Ser Ser Thr
1015 1020 1025

Ile Ser Cys Asp Ser Pro Leu Glu Leu Gln Glu Glu Pro Gln
1030 1035

35 Gln Ala Glu Pro Glu Ala Gln Leu Glu Gln Pro Gln Asp Ser
1040 1045 1050

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Gly Cys Pro Gly Pro Leu Ala Glu Ala Glu Asp Ser Phe Leu
1055 1060 1065

5 Glu Gln Pro Gln Asp Ser Gly Cys Pro Gly Pro Leu Ala Glu
1070 1075 1080

Ala Glu Asp Ser Phe Leu
1085

10 (2) INFORMATION FOR SEQ ID NO: 11:

(i) SEQUENCE CHARACTERISTICS:

15 (A) LENGTH: 16 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

20

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:

TCGACGCGCG ATG GAG

16

30

35

- 67 -

We claim:

1. An isolated DNA sequence encoding the Kinase insert Domain containing Receptor.

2. The DNA sequence of Claim 1 wherein said sequence is a human gene.

3. An isolated DNA sequence comprising a DNA sequence capable of hybridizing with a DNA sequence encoding the amino acid sequence of Figure 7.

4. A method for the production of a growth factor receptor which comprises transforming a host cell with the DNA sequence of Claim 3 and culturing the host cell under conditions which result in expression of the gene by an expression vector.

5. The method of Claim 4 wherein the host cell is a bacteria, virus, yeast, insect or mammalian cell line.

6. The method of Claim 5 wherein the host cell is a COS-1 cell, NIH3T3 fibroblast or CMT-3 monkey kidney cell.

7. The method of Claim 5 where the expression vector is pcDNA1tkpASP expression vector.

8. A lambda gt11 phage harboring the clone BTIII1081.8 (ATCC accession number 40,931) or the clone BTIII129.5 (ATCC Accession number 40,975).

9. A plasmid pBlueScript KS which contains the clone BTIV169 (ATCC accession number 75200).

10. An isolated growth factor receptor designated the Kinase insert Domain containing Receptor.

11. The receptor of Claim 10 comprising the amino acid sequence of Figure 7.

12. The receptor of Claim 10 encoded by an isolated DNA sequence comprising a DNA sequence capable

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of hybridizing with a DNA sequence encoding the amino acid sequence of Figure 7.

13. A biologically active protein fragment which retains the receptor activity of the receptor of Claim 10.

14. An isolated DNA sequence encoding a biologically active protein fragment which retains the receptor activity of an isolated growth factor receptor designated the Kinase insert Domain containing Receptor.

15. An oligonucleotide primer consisting of an oligonucleotide primer having 21 bases and having a sequence depicted for Primer 1 in Figure 2.

16. An oligonucleotide primer consisting of an oligonucleotide primer having 29 bases and having a sequence depicted for Primer 2 in Figure 2.

17. The 363 base pair product having the sequence depicted in Figure 4, or a biological equivalent of said sequence.

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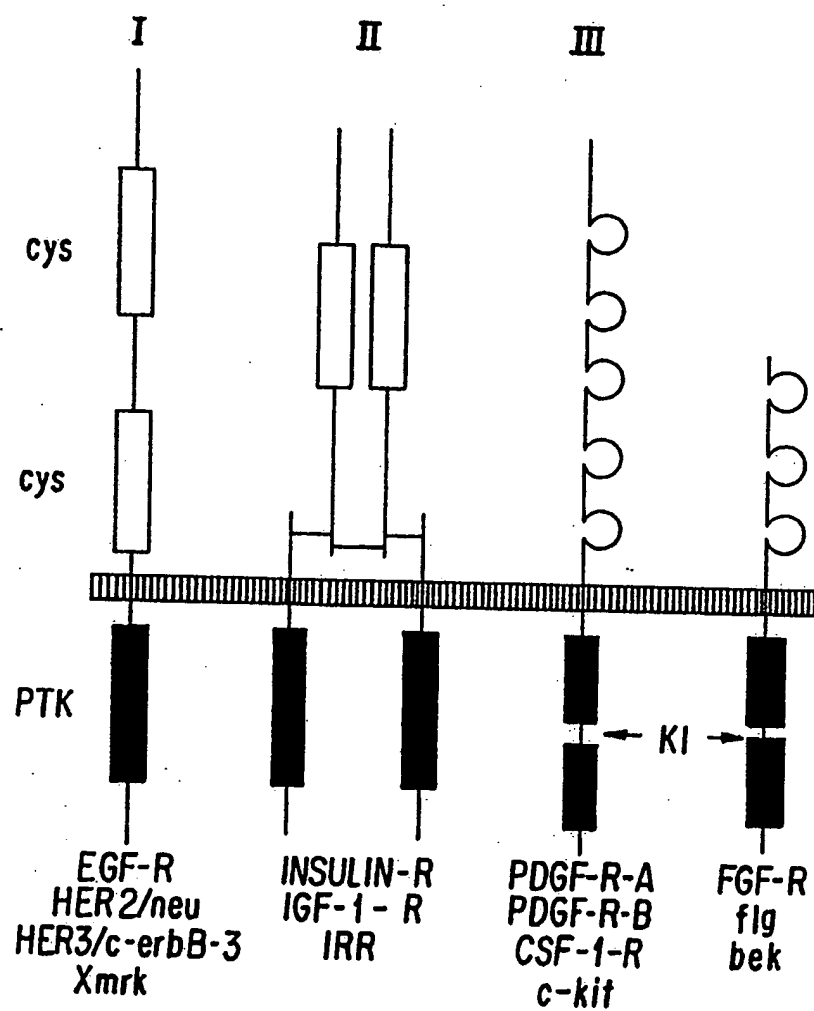


FIG. 1

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FIG. 2

PRIMER 1RECEPTOR

PDGF	AAC	CTG	TTG	GGG	GCC	TGC	ACC
ck1+	T	A	T	A			
CSF	T			A			
FGF		C					G

PRIMER 1

GTCGAC	AAC	CTG	TTG	GGG	GCC	TGC	AAC
	T			A			

PRIMER 2RECEPTOR

PDGF	CAC	AGA	GAC	CTG	GCG	GCT	AGG	AAC	GTG	CT
ck1+				T	GA	C	A	T		A
CSF		C	G	G	A	GC	C	T		
FGF		C			C	C			T	C

CONSENSUS

CAC	AGA	GAC	CTG	GCC	GCT	AGI	AAC	GTG	CT
	C			T	C			T	

PRIMER 2

GAATTC	AG	CAC	GTT	ICT	AGC	CGC	CAG	GTC	TCT	GTG
			T		G	T				G

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420bp

230bp

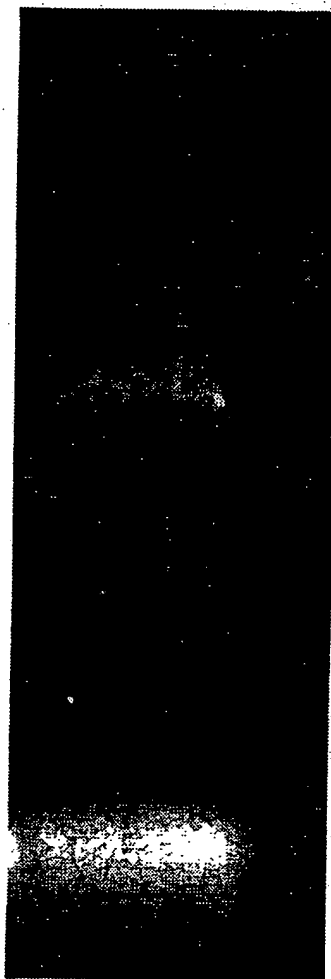


FIG. 3

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[illegible]

FIG. 4A

G	T	C	G	A	C	A	T	C	T	G	T	G	G	G	G	C	C	T	G	C	A	T	T	A	G	C	C	T	C
C	A	C	C	G	G	A	T	G	C	T	G	C	T	A	C	T	A	C	T	A	C	T	A	G	C	G	A	T	C
G	A	T	G	A	C	C	G	C	G	A	T	G	C	T	A	C	T	A	C	T	A	G	C	G	A	T	A	G	C
A	A	G	A	C	C	A	T	G	C	A	T	G	G	A	T	G	A	G	A	T	G	C	A	G	A	T	A	G	C
A	A	T	A	G	A	T	G	C	C	A	G	C	C	A	G	A	T	G	A	G	A	T	A	G	A	T	A	G	C

FIG. 4B

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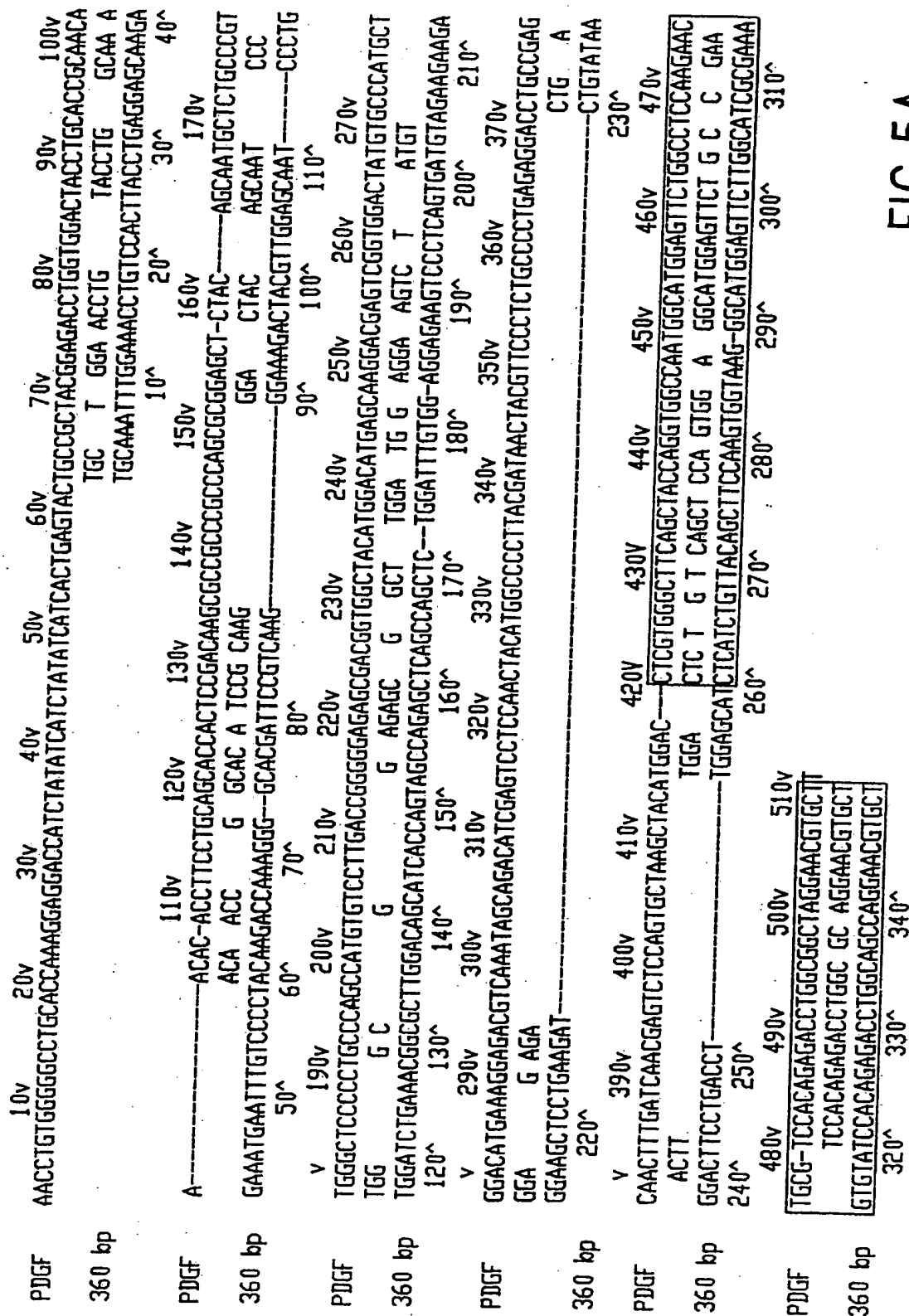


FIG. 5A

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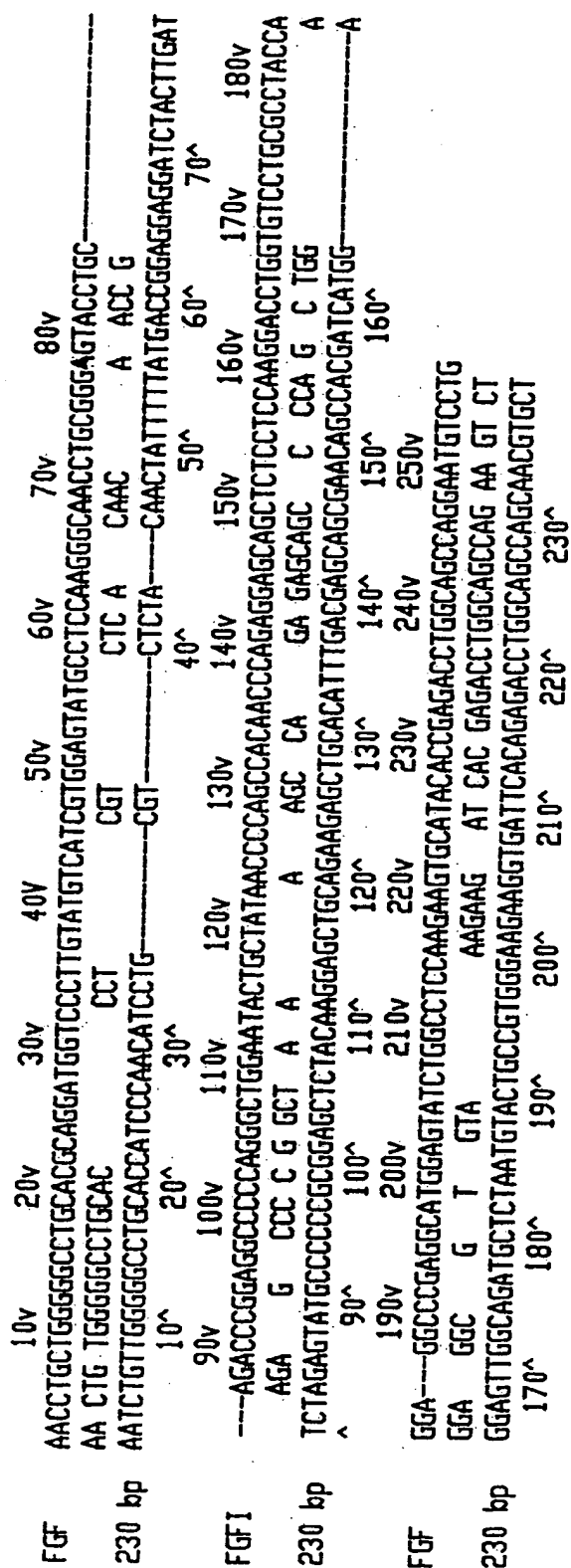


FIG. 5B

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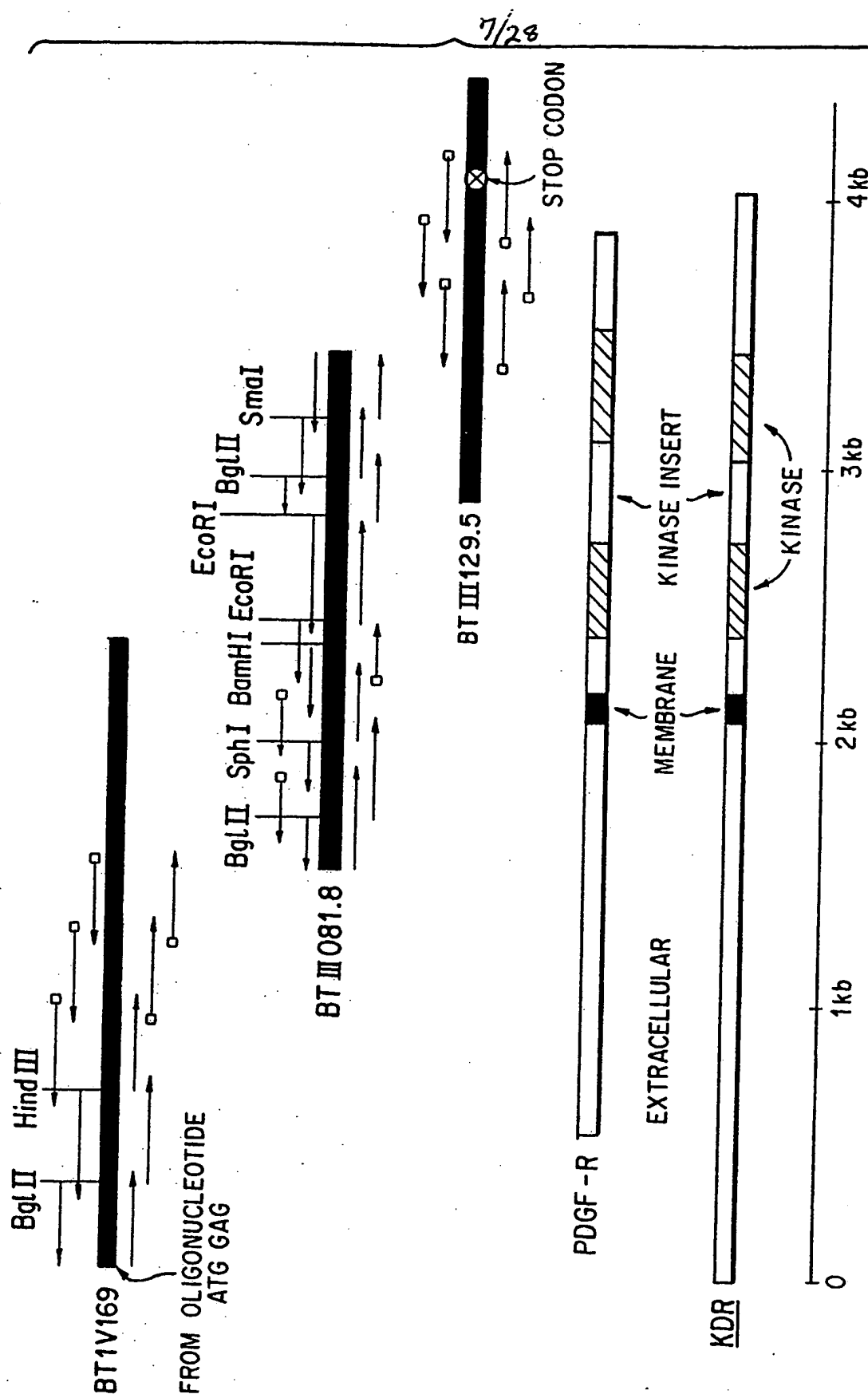


FIG. 6

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10 * 20 * 30 * 40 * 50 *
 ATG GAG AGC AAG GTG CTG CTG GCC GTC TGG CTC TGC GTG GAG ACC CCG
 Met Glu Ser Lys Val Leu Leu Ala Val Ala Leu Trp Leu Cys Val Glu Thr Arg>
 60 * 70 * 80 * 90 * 100 *
 GCC GCC TCT GTG GGT TTG CCT AGT GTT TCT CTT GAT CTG CCC AGG CTC AGC ATA
 Ala Ala Ser Val Gly Leu Pro Ser Val Ser Leu Asp Leu Pro Arg Leu Ser Ile>
 110 * 120 * 130 * 140 * 150 * 160 *
 CAA AAA GAC ATA CTT ACA ATT AAG GCT AAT ACA ACT CTT CAA ATT ACT TGC AGG
 Gln Lys Asp Ile Leu Thr Ile Lys Ala Asn Thr Thr Leu Gln Ile Thr Cys Arg>
 170 * 180 * 190 * 200 * 210 *
 GGA CAG AGG GAC TTG GAC TGG CTT TGG CCC AAT AAT CAG AGT GGC AGT GAG CAA
 Gly Gln Arg Asp Leu Asp Trp Leu Trp Pro Asn Asn Gln Ser Gly Ser Glu Gln>
 220 * 230 * 240 * 250 * 260 * 270 *
 AGG GTG GAG GTG ACT GAG TGC AGC GAT GGC CTC TTC TGT AAG ACA CTC ACA ATT
 Arg Val Glu Val Thr Glu Cys Ser Asp Gly Leu Phe Cys Lys Thr Leu Thr Ile>
 280 * 290 * 300 * 310 * 320 *
 CCA AAA GTG ATC GGA AAT GAC ACT GGA GCC TAC AAG TGC TTC TAC CGG GAA ACT
 Pro Lys Val Ile Gly Asn Asp Thr Gly Ala Tyr Lys Cys Phe Tyr Arg Glu Thr>

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FIG. 7A

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330 * 340 * 350 * 360 * 370 *
 GAC TTG GCC TCG GTC ATT TAT GTC TAT GTT CAA GAT TAC AGA TCT CCA TTT ATT
 Asp Leu Ala Ser Val Ile Tyr Val Tyr Val Gln Asp Tyr Arg Ser Pro Phe Ile>
 380 * 390 * 400 * 410 * 420 * 430 *
 GCT TCT GTT AGT GAC CAA CAT GGA GTC GTG TAC ATT ACT GAG AAC AAA AAC AAA
 Ala Ser Val Ser Asp Gln His Gly Val Val Tyr Ile Thr Glu Asn Lys Asn Lys>
 440 * 450 * 460 * 470 * 480 *
 ACT GTG GTG ATT CCA TGT CTC GGG TCC ATT TCA AAT CTC AAC GTG TCA CTT TGT
 Thr Val Val Ile Ile Pro [Cys] Leu Gly Ser Ile Ser Asn Leu Asn Val Ser Leu [Cys]>
 490 * 500 * 510 * 520 * 530 * 540 ^{9/24} *
 GCA AGA TAC CCA GAA AAG AGA TTT GTT CCT GAT GGT AAC AGA ATT TCC TGG GAC
 Ala Arg Tyr Pro Glu Lys Arg Phe Val Pro Asp Gly Asn Arg Ile Ser Trp Asp>
 550 * 560 * 570 * 580 * 590 *
 AGC AAG AAG GGC TTT ACT ATT CCC AGC TAC ATG ATC AGC TAT GCT GGC ATG GTC
 Ser Lys Lys Gly Phe Thr Ile Pro Ser Tyr Met Ile Ser Tyr Ala Gly Met Val>
 600 * 610 * 620 * 630 * 640 *
 TTC TGT GAA GCA AAA ATT AAT GAT GAA AGT TAC CAG TCT ATT ATG TAC ATA GTT
 Phe [Cys] Glu Ala Lys Ile Asn Asp Glu Ser Tyr Gln Ser Ile Met Tyr Ile Val>

FIG. 7B

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650 * 660 670 680 690 700
 GTC GTT GTA GGG TAT AGG ATT TAT GAT GTG GTT CTG AGT CCG TCT CAT GGA ATT
 Val Val Val Gly Tyr Arg Ile Tyr Asp Val Val Leu Ser Pro Ser His Gly Ile>
 710 * 720 730 740 750
 GAA CTA TCT GTT GGA GAA AAG CTT GTC TTA AAT TGT ACA GCA AGA ACT GAA CTA
 Glu Leu Ser Val Gly Glu Lys Lys Leu Val Leu Asn [Cys] Thr Ala Arg Thr Glu Leu>
 760 * 770 780 790 800 810
 AAT GTG GGG ATT GAC TTC AAC TGG GAA TAC CCT TCT TCG AAG CAT CAG CAT AAG
 Asn Val Gly Ile Asp Phe Asn Trp Glu Tyr Pro Ser Ser Lys His Gln His Lys>
 820 * 830 840 850 860
 AAA CTT GTA AAC CGA GAC CTA AAA ACC CAG TCT GGG AGT GAG ATG AAG AAA TTT
 Lys Leu Val Asn Arg Asp Leu Lys Thr Gln Ser Gly Ser Glu Met Lys Lys Phe>
 870 * 880 890 900 910
 TTG AGC ACC TTA ACT ATA GAT GGT GTA ACC CGG AGT GAC CAA GGA TTG TAC ACC
 Leu Ser Thr Leu Thr Ile Asp Gly Val Thr Arg Ser Asp Gln Gly Leu Tyr Thr>
 920 * 930 940 950 960 970
 TGT GCA GCA TCC AGT GGG CTG ATG ACC AAG AAG AAC AGC ACA TTT GTC AGG GTC
 [Cys] Ala Ala Ser Ser Gly Leu Met Thr Lys Lys Asn Ser Thr Phe Val Arg Val>

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FIG. 7C

980	990	1000	1010	1020
CAT GAA AAA CCT TTT GTT GCT TTT GGA AGT GGC ATG GAA TCT CTG GTG GAA GCC	*	*	*	*
His Glu Lys Pro Phe Val Ala Phe Gly Ser Gly Met Glu Ser Leu Val Glu Ala>				
1030	1040	1050	1060	1070
ACG GTG GGG GAG CGT GTC AGA ATC CCT GCG AAG TAC CTT GGT TAC CCA CCC CCA	*	*	*	*
Thr Val Gly Glu Arg Val Arg Ile Pro Ala Lys Tyr Leu Gly Tyr Pro Pro Pro>				
1090	1100	1110	1120	1130
GAA ATA AAA TGG TAT AAA AAT GGA ATA CCC CTT GAG TCC AAT CAC ACA ATT AAA	*	*	*	*
Glu Ile Lys Lys Trp Tyr Lys Asn Gly Ile Pro Leu Glu Ser Asn His Thr Ile Lys>				
1140	1150	1160	1170	1180
CGG GGG CAT GTA CTG ACG ATT ATG GAA GTG AGT GAA AGA GAC ACA GGA AAT TAC	*	*	*	*
Ala Gly His Val Leu Thr Ile Met Glu Val Ser Glu Arg Asp Thr Gly Asn Tyr>				
1190	1200	1210	1220	1230
ACT GTC ATC CTT ACC AAT CCC ATT TCA AAG GAG AAG CAG AGC CAT GTG GTC TCT	*	*	*	*
Thr Val Ile Leu Thr Asn Pro Ile Ser Lys Glu Lys Gln Ser His Val Val Ser>				
1250	1260	1270	1280	1290
CTG GTT GTG TAT GTC CCA CCC CAG ATT GGT GAG AAA TCT CTA ATC TCT CCT GTG	*	*	*	*
Leu Val Val Tyr Val Pro Pro Gln Ile Gly Glu Lys Ser Leu Ile Ser Pro Val>				

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FIG. 7D

1300 * 1310 * 1320 * 1330 * 1340 * 1350 *
 GAT TCC TAC CAG TAC GGC ACC ACT CAA ACG CTG ACA TGT ACG GTC TAT GCC ATT
 Asp Ser Tyr Gln Tyr Gly Thr Thr Gln Thr Leu Thr Cys Thr Val Tyr Ala Ile>

 1360 * 1370 * 1380 * 1390 * 1400 *
 CCT CCC CCG CAT CAC ATC CAC TGG TAT TGG CAG TTG GAG GAA GAG TGC GCC AAC
 Pro Pro Pro His His Ile His Trp Tyr Trp Gln Leu Glu Glu Cys Ala Asn>

 1410 * 1420 * 1430 * 1440 * 1450 *
 GAG CCC AGC CAA GCT GTC TCA GTG ACA AAC CCA TAC CCT TGT GAA GAA TGG AGA
 Glu Pro Ser Gln Ala Val Ser Val Thr Asn Pro Tyr Pro Cys Glu Glu Trp Arg>

 1460 * 1470 * 1480 * 1490 * 1500 * 1510 *
 AGT GTG GAG GAC TTC CAG GGA GGA AAT AAA ATT GAA GTT AAT AAA AAT CAA TTT
 Ser Val Glu Asp Phe Gln Gly Gly Asn Lys Ile Glu Val Asn Lys Asn Gln Phe>

 1520 * 1530 * 1540 * 1550 * 1560 *
 GCT CTA ATT GAA GGA AAA AAC AAA ACT GTA AGT ACC CTT GTT ATC CAA GCG GCA
 Ala Leu Ile Glu Gly Lys Lys Asn Lys Thr Val Ser Thr Thr Leu Val Ile Gln Ala Ala>

 1570 * 1580 * 1590 * 1600 * 1610 * 1620 *
 AAT GTG TCA GCT TTG TAC AAA TGT GAA GCG GTC AAC AAA GTC GGG AGA GGA GAG
 Asn Val Ser Ala Leu Tyr Lys Cys Glu Ala Val Asn Lys Val Gly Arg Gly Glu>

FIG. 7E

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1630	1640	1650	1660	1670
AGG GTG ATC TCC TTC CAC GTG ACC AGG GGT CCT GAA ATT ACT TTG CAA CCT GAC Arg Val Ile Ser Phe His Val Thr Arg Gly Pro Glu Ile Thr Leu Gln Pro Asp>				
1680	1690	1700	1710	1720
ATG CAG CCC ACT GAG CAG GAG AGC GTG TCT TTG TGG TGC ACT GCA GAC AGA TCT Met Gln Pro Thr Glu Gln Glu Ser Val Ser Leu Trp <u>Cys</u> Thr Ala Asp Arg Ser>				
1730	1740	1750	1760	1770
ACG TTT GAG AAC CTC ACA TGG TAC AAG CTT GGC CCA CAG CCT CTG CCA ATC CAT Thr Phe Glu Asn Leu Thr Thr Trp Tyr Lys Leu Gly Pro Gln Pro Leu Pro Ile His>				
1790	1800	1810	1820	1830
GTG GGA GAG TTG CCC ACA CCT GTT TGC AAG AAC TTG GAT ACT CTT TGG AAA TTG Val Gly Glu Leu Pro Thr Thr Pro Val <u>Cys</u> Lys Asn Leu Asp Thr Leu Trp Lys Leu>				
1840	1850	1860	1870	1880
AAT GCC ACC ATG TTC TCT AAT AGC ACA AAT GAC ATT TTG ATC ATG GAG CTT AAG Asn Ala Thr Met Phe Ser Asn Ser Thr Asn Asp Ile Leu Ile Met Glu Leu Lys>				
1900	1910	1920	1930	1940
AAT GCA TCC TTG CAG GAC CAA GGA GAC TAT GTC TGC CTT GCT CAA GAC AGG AAG Asn Ala Ser Leu Gln Asp Gln Gly Asp Tyr Val Cys Leu Ala Gln Asp Arg Lys>				

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FIG. 7F

1950	1960	1970	1980	1990
* ACC AAG AAA AGA CAT TGC GTG GTC AGG CAG CTC ACA GAG CGT GTG GCA Thr Lys Lys Arg His <u>Cys</u> Val Val Arg Gln Leu Thr Val Leu Glu Arg Val Ala>				
2000	2010	2020	2030	2040
* CCC ACG ATC ACA GGA AAC CTG GAG AAT CAG ACG ACA AGT ATT GGG GAA AGC ATC Pro Thr Ile Thr Gly Asn Leu Glu Asn Gln Thr Thr Ser Ile Gly Glu Ser Ile>				
2060	2070	2080	2090	2100
* GAA GTC TCA TGC ACG GCA TCT GGG AAT CCC CCT CCA CAG ATC ATG TGG TTT AAA Glu Val Ser <u>Cys</u> Thr Ala Ser Gly Asn Pro Pro Pro Gln Ile Met Trp Phe Lys>				
2110	2120	2130	2140	2150
* GAT AAT GAG ACC CTT GTA GAA GAC TCA GGC ATT GTA TTG AAG GAT GGG AAC CCG Asp Asn Glu Thr Leu Val Glu Asp Ser Gly Ile Val Leu Lys Asp Gly Asn Arg>				
2170	2180	2190	2200	2210
* AAC CTC ACT ATC CGC AGA GTG AGG AAG GAG GAC GAA GGC CTC TAC ACC TGC CAG Asn Leu Thr Ile Arg Arg Val Arg Lys Glu Asp Glu Gly Leu Tyr Thr <u>Cys</u> Gln>				
2220	2230	2240	2250	2260
* GCA TGC AGT GTT CTT GGC TGT GCA AAA GTG GAG GCA TTT TTC ATA ATA GAA GGT Ala <u>Cys</u> Ser Val Leu Gly <u>Cys</u> Ala Lys Val Glu Ala Phe Phe Ile Ile Glu Gly>				

FIG. 7G

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2600	2610	2620	2630	2640
GTA GCA GTC AAA ATG TTG AAA GAA GGA GCA ACA CAC AGT GAG CAT CGA GCT CTC	*	*	*	*
Val Ala Val Lys Met Leu Lys Glu Gly Ala Thr His Ser Glu His Arg Ala Leu>				
2650	2660	2670	2680	2690
ATG TCT GAA CTC AAG ATC CTC ATT CAT ATT GGT CAC CAT CTC AAT GTG GTC AAC	*	*	*	*
Met Ser Glu Leu Lys Ile Leu Ile His Ile Gly His Leu Asn Val Val Asn>				
2710	2720	2730	2740	2750
CTT CTA GGT GCC TGT ACC AAG CCA GGA GGG CCA CTC ATG GTG ATT GTG GAA TTC	*	*	*	*
Leu Leu Gly Ala Cys Thr Lys Lys Pro Gly Gly Pro Leu Met Val Ile Val Glu Phe>				
2760	2770	2780	2790	2800
TGC AAA TTT GGA AAC CTG TCC ACT TAC TAC CTG AGG AGC AAG AGA AAT GAA TTT GTC	*	*	*	*
Cys Lys Phe Gly Asn Leu Ser Thr Tyr Leu Arg Ser Lys Arg Asn Glu Phe Val>				
2810	2820	2830	2840	2850
CCC TAC AAG ACC AAA GGG GCA CGA TTC CGT CAA GGG AAA GAC TAC GTT GGA GCA	*	*	*	*
Pro Tyr Lys Thr Lys Gly Ala Arg Phe Arg Gln Gly Lys Asp Tyr Val Gly Ala>				
2870	2880	2890	2900	2910
ATC CCT GTG GAT CTG AAA CGG CGC TTG GAC AGC ATC ACC AGT AGC CAG AGC TCA	*	*	*	*
Ile Pro Val Asp Leu Lys Arg Arg Leu Asp Ser Ile Thr Ser Ser Gln Ser Ser>				

FIG. 7I

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3250 * 3260 * 3270 * 3280 * 3290 *
 GTG TAC ACA ATC CAG AGT GAC GTC TGG TCT TTT GGT GTT TTG CTG TGG GAA ATA
 Val Tyr Thr Ile Gln Ser Asp Val Trp Ser Phe Gly Val Leu Leu Trp Glu Ile>
 3300 * 3310 * 3320 * 3330 * 3340 *
 TTT TCC TTA GGT GCT TCT CCA TAT CCT GGG GTA AAG ATT GAT GAA GAA TTT TGT
 Phe Ser Leu Gly Ala Ser Pro Tyr Pro Gly Val Lys Ile Asp Glu Glu Phe Cys>
 3350 * 3360 * 3370 * 3380 * 3390 * 3400 *
 AGG CGA TTG AAA GAA GGA ACT AGA ATG AGG GCC CCT GAT TAT ACT ACA CCA GAA
 Arg Arg Leu Lys Glu Gly Thr Arg Met Arg Ala Pro Asp Tyr Thr Thr Pro Glu>
 3410 * 3420 * 3430 * 3440 * 3450 *
 ATG TAC CAG ACC ATG CTG GAC TGC TGG CAC GGG GAG CCC AGT CAG AGA CCC ACG
 Met Tyr Gln Thr Met Leu Asp Cys Trp His Gly Glu Pro Ser Gln Arg Pro Thr>
 3460 * 3470 * 3480 * 3490 * 3500 * 3510 *
 TTT TCA GAG TTG GTG GAA CAT TTG GGA AAT CTC TTG CAA GCT AAT GCT CAG CAG
 Phe Ser Glu Leu Val Glu His Leu Gly Asn Leu Leu Gln Ala Asn Ala Gln Gln>
 3520 * 3530 * 3540 * 3550 * 3560 *
 GAT GGC AAA GAC TAC ATT GTT CTT CCG ATA TCA GAG ACT TTG AGC ATG GAA GAG
 Asp Gly Lys Asp Tyr Ile Val Leu Pro Ile Ser Glu Thr Leu Ser Met Glu Glu>

FIG. 7K

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3570	3580	3590	3600	3610
* GAT TCT GGA CTC TCT CTG CCT ACC TCA CCT GTT TCC TGT ATG GAG GAG GAG GAA	*	*	*	*
Asp Ser Gly Leu Ser Leu Pro Thr Ser Pro Val Ser Cys Met Glu Glu Glu>				
3620	3630	3640	3650	3660
* GTA TGT GAC CCC AAA TTC CAT TAT GAC AAC ACA GCA GGA ATC AGT CAG TAT CTG	*	*	*	*
Val Cys Asp Pro Lys Phe His Tyr Asp Asn Thr Ala Gly Ile Ser Gln Tyr Leu>				
3680	3690	3700	3710	3720
* CAG AAC AGT AAG CGA AAG AGC CGG CCT GTG AGT GTA AAA ACA TTT GAA GAT ATC	*	*	*	*
Gln Asn Ser Lys Arg Lys Ser Arg Pro Val Ser Val Lys Thr Phe Glu Asp Ile>				
3730	3740	3750	3760	3770
* CCG TTA GAA GAA CCA GAA GTA AAA GTA ATC CCA GAT GAC AAC CAG ACG GAC AGT	*	*	*	*
Pro Leu Glu Glu Pro Glu val Lys Val Ile Pro Asp Asp Asn Gln Thr Asp Ser>				
3790	3800	3810	3820	3830
* GGT ATG GTT GCT GCC TCA GAA GAG CTG AAA ACT TTG GAA GAC AGA ACC AAA TTA	*	*	*	*
Gly Met Val Leu Ala Ser Glu Glu Leu Lys Thr Leu Glu Asp Arg Thr Lys Leu>				
3840	3850	3860	3870	3880
* TCT CCA TCT TTT GGT GGA ATG GTG CCC AGC AAA AGC AGG GAG TCT GTG GCA TCT	*	*	*	*
Ser Pro Ser Phe Gly Gly Met val Pro Ser Lys Ser Arg Glu Ser Val Ala Ser>				

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FIG. 7L

3890 * 3900 * 3910 * 3920 * 3930 * 3940 *
 GAA GGC TCA AAC CAG ACA AGC GGC TAC CAG TCC GGA TAT CAC TCC GAT GAC ACA
 Glu Gly Ser Asn Gln Thr Ser Gly Tyr Gln Ser Gly Tyr His Ser Asp Thr>
 3950 * 3960 * 3970 * 3980 * 3990 *
 GAC ACC ACC GTG TAC TCC AGT GAG GAA GCA GAA CTT TTA AAG CTG ATA GAG ATT
 Asp Thr Thr Val Tyr Ser Ser Glu Glu Ala Glu Leu Lys Leu Ile Glu Ile>
 4000 * 4010 * 4020 * 4030 * 4040 * 4050 *
 GGA GTG CAA ACC GGT AGC ACA GCC CAG ATT CTC CAG CCT GAC ACG GGG ACC ACA
 Gly Val Gln Thr Gly Ser Thr Ala Gln Ile Leu Gln Pro Asp Thr Gly Thr Thr>
 4060 * 4070 *
 CTG AGC TCT CCT CCT GTT TAA
 Leu Ser Ser Pro Pro Val ***

FIG. 7M

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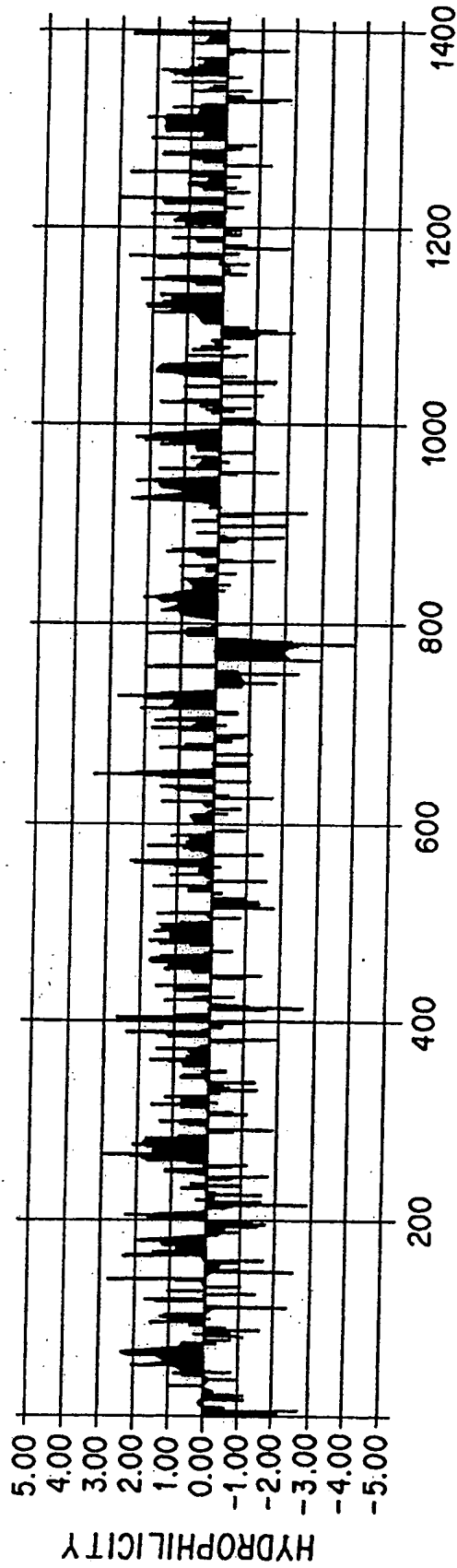


FIG. 8

KDR 787 GTVKRANGGELKTGYLSIVMDPDELPLDEHCERLPYDASKWEFPRDRNLGK
 ckit 543 L***YLQKPMYEVQWKVVEEINGNNYVYIDPTQ***H-*****N**SF**
 CSF1 536 LLY*YKQPKYQVRWKIIESYEGNSYTFIDPTQ***NE-*****NN*QF**
 PDGF 522 MLWQKKPRYEIRWKVIESVSSDGHYIYVDPVQ***-ST*****QLV**R

* * *

KDR 839 PLGRGAFGQIEADAFGIDKTATCRTVAVKMLKEGATHSEHRAHMLSELKILI
 ckit 594 T**A***KVVAEY*LI*SDAAM*****PS*HLT*RE*****V*S
 CSF1 587 T**A***KVV**T***LG*EDAVLK*****ST*HAD*KE*****MS
 PDGF 573 T**S***V**T**H*LSHSQATMK*****ST*RSS*KQS*****MS

KDR 891 HIGHHLNVVNLGACTKPGGPLMVIVEFCKFGNLSYLRSKRNEFVPYKTKG
 ckit 646 YL*N*M*I*****I-***TL**T*Y*CY*D*LNF**R**DS*ICS*QED
 CSF1 639 *L*Q*E*I*****H-***VL**T*Y*CY*D*LNF**R*AEAMLGPSLSP
 PDGF 625 *L*P*****I*YI*T*Y*RY*D*VD**HRNKHT*LQRHSNK

KDR 943 ARFRQKDYVGAIPVDLKRRLDSIT-SSQSSASSGFVEEKS-----SDV
 ckit 697 HAEA-A-L*KNLLHSKESSCS-DS*N-E-----YMDMKPGVS--YVPT--KA
 CSF1 690 QODPE*GVDYKN*HLEK*YVRRDSGF**GVDTYVEMRPVSTSS-NDSE*EQ
 PDGF 676 HCPPSAEL*SN*LP*GFSLP SHLNLTGESDGGYMDMSKDESIDYVPMMLDMKG

KDR 987 EEEEAPELKYKF-----LTLEHLICYSFQV
 ckit 737 D-KRRSVRIGSYI-----ERDVTPAIMEDELA*D**D*LSF*Y**
 CSF1 741 DLDKEDGRPL-----E*RD*LHF*S**
 PDGF 728 DIKY*DIESPSYMAPYDNYVPSAPERTYRATLINDSPV-*SYTD*VGF*Y**

KDR 1013 AKGMEFLASRKC IHRDLAARNILLSEKNVVKICDFGLARDIYKDPDYVRKGD

FIG. 9A

SUBSTITUTE SHEET

IDENTIFICATION OF *kdp* mRNA

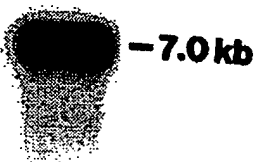


FIG. 10

**IDENTIFICATION OF *kdp* GENE
BY SOUTHERN ANALYSIS**



1 2 3 4

FIG. 11

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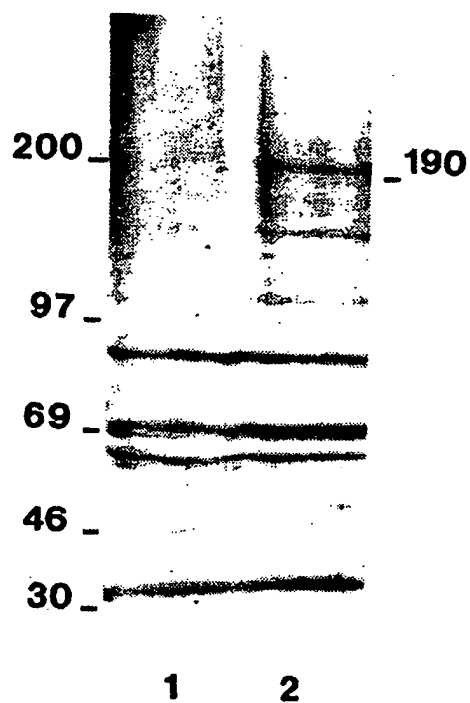


FIG. 12

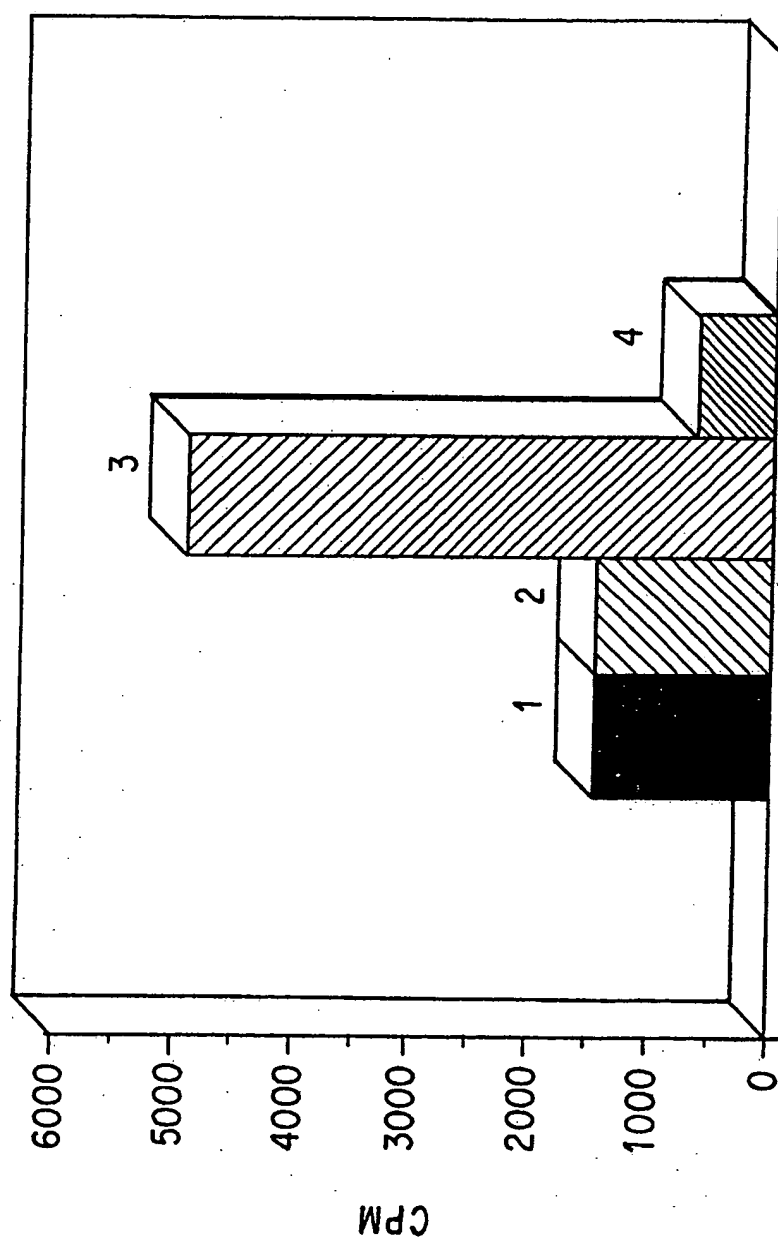


FIG. 13

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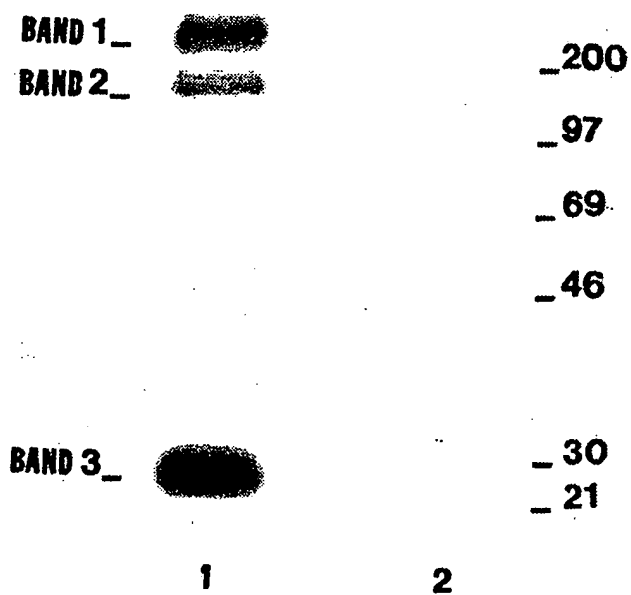


FIG. 14

INTERNATIONAL SEARCH REPORT

International Application No. PCT/US92/01300

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) ³ According to International Patent Classification (IPC) or to both National Classification and IPC IPC (5): C07K 3/00, 13/00; C07H 21/00; C12P 21/06, 21/02, 21/04; C12N 15/00 US CL : 530/387; 536/27; 435/69.1, 70.1, 71.1, 320.1																	
II. FIELDS SEARCHED <div style="text-align: center; border: 1px solid black; padding: 2px;">Minimum Documentation Searched⁴</div> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 20%;">Classification System</th> <th>Classification Symbols</th> </tr> <tr> <td style="text-align: center; vertical-align: top;">U.S.</td> <td>530/387; 536/27; 435/69.1, 70.1, 71.1, 320.1</td> </tr> </table> <div style="text-align: center; border: 1px solid black; padding: 2px;">Documentation Searched other than Minimum Documentation to the extent that such Documents are included in the Fields Searched⁵</div> <p>APS, DIALOG search terms: type III receptor tyrosine kinase</p>			Classification System	Classification Symbols	U.S.	530/387; 536/27; 435/69.1, 70.1, 71.1, 320.1											
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III. DOCUMENTS CONSIDERED TO BE RELEVANT ¹⁴ <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 10%;">Category¹⁵</th> <th style="width: 60%;">Citation of Document,¹⁶ with indication, where appropriate, of the relevant passages¹⁷</th> <th style="width: 30%;">Relevant to Claim No. ¹⁸</th> </tr> </thead> <tbody> <tr> <td style="vertical-align: top;">Y,P</td> <td>Proc. Natl. Acad. Sci., Volume 88, Issued 1991, W. Mathews et al., "A receptor tyrosine kinase cDNA isolated from a population of enriched primitive hematopoietic cells and exhibiting close genetic linkage to c-kit", pages 9026-9030, see entire document.</td> <td style="vertical-align: top;">1-17</td> </tr> <tr> <td style="vertical-align: top;">Y</td> <td>Proc. Natl. Acad. Sci., Volume 86, Issued March 1989, A.F. Wilks, "Two putative protein-tyrosine kinases identified by application of the polymerase chain reaction", pages 1603-1607, see entire document.</td> <td style="vertical-align: top;">1-17</td> </tr> <tr> <td style="vertical-align: top;">X,P</td> <td>Oncogene, Volume 6, issued 1991, B.I. Terman et al., "Identification of a new endothelial cell growth factor receptor tyrosine kinase", pages 1677-1683, see entire document.</td> <td style="vertical-align: top;">1-17</td> </tr> <tr> <td style="vertical-align: top;">A</td> <td>Oncogene, volume 3, issued 1988, M. Ruta et al., "A novel protein tyrosine kinase gene whose expression is modulated during endothelial cell differentiation", pages 9-15, see entire document.</td> <td style="vertical-align: top;">1-17</td> </tr> </tbody> </table>			Category ¹⁵	Citation of Document, ¹⁶ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸	Y,P	Proc. Natl. Acad. Sci., Volume 88, Issued 1991, W. Mathews et al., "A receptor tyrosine kinase cDNA isolated from a population of enriched primitive hematopoietic cells and exhibiting close genetic linkage to c-kit", pages 9026-9030, see entire document.	1-17	Y	Proc. Natl. Acad. Sci., Volume 86, Issued March 1989, A.F. Wilks, "Two putative protein-tyrosine kinases identified by application of the polymerase chain reaction", pages 1603-1607, see entire document.	1-17	X,P	Oncogene, Volume 6, issued 1991, B.I. Terman et al., "Identification of a new endothelial cell growth factor receptor tyrosine kinase", pages 1677-1683, see entire document.	1-17	A	Oncogene, volume 3, issued 1988, M. Ruta et al., "A novel protein tyrosine kinase gene whose expression is modulated during endothelial cell differentiation", pages 9-15, see entire document.	1-17
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<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>¹⁵ Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 50%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"Z" document member of the same patent family</p> </div> </div>																	
IV. CERTIFICATION <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; padding: 5px;"> Date of the Actual Completion of the International Search² <div style="text-align: center; font-weight: bold;">13 MAY 1992</div> </td> <td style="width: 50%; padding: 5px;"> Date of Mailing of this International Search Report² <div style="text-align: center; font-weight: bold;">19 MAY 1992</div> </td> </tr> <tr> <td style="width: 50%; padding: 5px;"> International Searching Authority¹ <div style="text-align: center; font-weight: bold;">ISA/US</div> </td> <td style="width: 50%; padding: 5px;"> Signature of Authorized Officer¹⁹ <div style="text-align: center;"> Lorraine M. Spector, Ph.D. </div> </td> </tr> </table>			Date of the Actual Completion of the International Search ² <div style="text-align: center; font-weight: bold;">13 MAY 1992</div>	Date of Mailing of this International Search Report ² <div style="text-align: center; font-weight: bold;">19 MAY 1992</div>	International Searching Authority ¹ <div style="text-align: center; font-weight: bold;">ISA/US</div>	Signature of Authorized Officer ¹⁹ <div style="text-align: center;"> Lorraine M. Spector, Ph.D. </div>											
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FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

A Y	Oncogene, volume 5, issued 1990, M. Shibuya et al., "Nucleotide sequence and expression of a novel human receptor-type tyrosine kinase gene (flt) closely related to the fms family", pages 519-524, see entire document.	1-17 17
Y	Proc. Natl. Acad. Sci., Volume 85, Issued May 1988, R.G.K. Gronwald et al., "Cloning and expression of a cDNA coding for the human platelet-derived growth factor receptor: Evidence for more than one receptor class", pages 3435-3439, see entire document.	15

V. ☐ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE¹

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☐ Claim numbers , because they relate to subject matter (1) not required to be searched by this Authority, namely:

2. ☐ Claim numbers , because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out (1), specifically:

3. ☐ Claim numbers , because they are dependent claims not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. ☒ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING²

This international Searching Authority found multiple inventions in this international application as follows:

- I. Claims 1-9 and 14-17, drawn to nucleic acids and expression thereof. Class 536, subclass 27 and Class 435, subclass 69.1.
- II. Claims 10-13, drawn to an isolated growth factor receptor. Class 530, subclass 387.

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application. (Telephone Practice)
2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:

3. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the international search authority did not invite payment of any additional fee.

Remark on protest

- ☐ The additional search fees were accompanied by applicant's protest.
☐ No protest accompanied the payment of additional search fees.

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category ¹⁶	Citation of Document, ¹⁶ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸
Y	Proc. Natl. Acad. Sci., Volume 86, Issued November 1989, M. Streuli et al., "A family of receptor-linked protein tyrosine phosphatases in humans and Drosophila", pages 8698-8702, see entire document.	1-14
Y	M.A. Innes et al., PCR Protocols, a guide to methods and applications, published 1990 by Academic Press (N.Y.), see page 10.	15, 16

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